

OM of: US-09-439-311-2 to: Issued\_Patents\_NA: \* out\_format : pfs  
 Date: Apr 17, 2002 3:10 AM  
 About: Results were produced by the GenCore software, version 4.5,  
 Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

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-MODEL:frame+P2n_model -DEV=xLP
-0-/gn2_1/USPTO_spool/US09439311/runat_16042002_134010_11703/app_query.fasta_1.395
-DB=Issued_Patents_NA -QFMT=fasta -SUFFIX=rni -GAPOP=12,000
-GAPEXT=4,000 -MINMATCH=0,100 -NOOPC=0,000 -LOOPCEN=0,000
-OGAPOP=4,500 -QGAPEXT=0,050 -XGAPOP=10,000 -XGAPEXT=0,500
-FGAPOP=6,000 -FGAPEXT=7,000 -YCAPOP=10,000 -YGAPEXT=0,500
-DELOP=6,000 -DELEXT=7,000 -START=1 -MATRIX=blosum62
-TRANS=human40_csl_LIST=45 -DOALIGN=0 -THR_SCORE=pct
-THR_MAX=100 -THR_MIN=0 -ALIGNM=15 -MODE=LOCAL -OUTFMT=pfs
-NORM=ext -MINLEN=0 -MAXLEN=0 -USER=US09439311@CSNL1_77
-NCPU=6 -ICPU=3 -LONGLOG -NO_XLPY -WAIT -THREADS=1
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Search information block:

Query: US-09-439-311-2

Database: Issued\_Patents\_NA: \*

Database sequences: 351203

Database length: 11323999

Search time (sec): 85.70000

score\_list:

Sequence      Strd Orig      Zscore      Escore Len      Documentation

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/cgn2_6/ptodata/2/1na/6B_COMB.seq:US-09-358-972-102 - 50.00 116.12 80.95 30 1
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/cgn2_6/ptodata/2/1na/6B_COMB.seq:US-09-358-972-103 + 50.00 116.12 80.95 30 1
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/cgn2_6/ptodata/2/1na/6B_COMB.seq:US-09-406-147-32 - 50.00 116.12 80.95 30 1
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/cgn2_6/ptodata/2/1na/6B_COMB.seq:US-09-406-147-34 + 50.00 116.12 80.95 30 1
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/cgn2_6/ptodata/2/1na/6A_COMBO.seq:US-09-081-180-29 + 43.00 96.03 1.1e+03 51 1
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/cgn2_6/ptodata/2/1na/6A_COMBO.seq:US-09-412-335-22 + 43.00 96.03 1.1e+03 51 1
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/cgn2_6/ptodata/2/1na/6A_COMBO.seq:US-09-219-012-80 + 40.00 88.19 2.9e+03 60 1
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/cgn2_6/ptodata/2/1na/6A_COMBO.seq:US-08-714-017-97 - 39.00 89.11 2.6e+03 47 1
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/cgn2_6/ptodata/2/1na/6B_COMBO.seq:US-08-475-609-97 - 39.00 89.11 2.6e+03 47 1
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/cgn2_6/ptodata/2/1na/5A_COMBO.seq:US-08-493-533-657-3 - 38.00 85.71 4.0e+03 53 1
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/cgn2_6/ptodata/2/1na/5B_COMBO.seq:US-08-720-4208-97 - 38.00 85.71 4.0e+03 53 1
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/cgn2_6/ptodata/2/1na/6A_COMBO.seq:US-08-813-507-82 - 37.00 86.79 3.5e+03 41 1
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/cgn2_6/ptodata/2/1na/6B_COMBO.seq:US-08-813-507-82 - 37.00 86.79 3.5e+03 41 1
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/cgn2_6/ptodata/2/1na/5A_COMBO.seq:US-08-975-699-4 - 37.00 84.50 4.6e+03 49 1
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/cgn2_6/ptodata/2/1na/5B_COMBO.seq:US-08-975-699-4 - 37.00 82.27 6.2e+03 60 1
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/cgn2_6/ptodata/2/1na/6B_COMBO.seq:US-08-977-089-4 - 37.00 82.27 6.2e+03 60 1
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/cgn2_6/ptodata/2/1na/5A_COMBO.seq:US-09-133-717-5 - 37.00 82.27 6.2e+03 60 1
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/cgn2_6/ptodata/2/1na/6B_COMBO.seq:US-09-133-717-5 - 37.00 82.27 6.2e+03 60 1
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/cgn2_6/ptodata/2/1na/5B_COMBO.seq:US-08-564-955-54 + 36.00 81.77 6.6e+03 53 1
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/cgn2_6/ptodata/2/1na/6B_COMBO.seq:US-09-358-972-102
/cgn2_6/ptodata/2/1na/6B_COMBO.seq:US-09-099-015-54 +
36.00 81.77 6.6e+03 53 1
/cgn2_6/ptodata/2/1na/6B_COMBO.seq:US-09-232-863-54 +
36.00 81.77 6.6e+03 53 1
/cgn2_6/ptodata/2/1na/6B_COMBO.seq:US-09-133-508A-54 +
36.00 81.77 6.6e+03 53 1
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seq\_documentation\_block:

Sequence 102, Application US/09358972

Patent No. 6235480

GENERAL INFORMATION:

APPLICANT: Shultz, John W

APPLICANT: Lewis, Martin K

APPLICANT: Lieppe, Donna

APPLICANT: Mandrekar, Michelle

APPLICANT: Kephart, Daniel B

APPLICANT: Rhodes, Richard B

APPLICANT: Olson, Ryan J

APPLICANT: Wood, Keith W

APPLICANT: Welch, Roy

TITLE OF INVENTION: Nucleic Acid Detection

FILE REFERENCE: PTO 103 6868/7528

CURRENT APPLICATION NUMBER: US/09-558,972

EARLIER APPLICATION NUMBER: 09/252,436

EARLIER FILING DATE: 1999-02-18

EARLIER APPLICATION NUMBER: 09/042,287

EARLIER FILING DATE: 1998-03-13

NUMBER OF SEQ ID NOS: 290

SEQ ID NO 102

LENGTH: 30

TYPE: DNA

ORGANISM: Campylobacter jejuni

FEATURE: OTHER INFORMATION: probe to Campylobacter jejuni

US-09-358-972-102

Alignment\_scores:

Quality: 50.00

Length: 10

Ratio: 5.000

Gaps: 0

Percent Identity: 100.000

Percent Similarity: 100.000

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alignment-block:

US-09-439-311-2 x US-09-358-972-102/rev ..

Align seg 1/1 to reverse of: US-09-358-972-102 from: 1 to: 30

align seg 1/1 to reverse of: US-09-358-972-102 from: 1 to: 30

align seg 1/1 to reverse of: US-09-358-972-102 from: 1 to: 30

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align seg 1/1 to reverse of: US-09-358-972-102 from: 1 to: 30

seq\_name: /cgn2\_6/ptodata/2/1na/6B\_COMBO.seq:US-09-358-972-103

seq\_documentation\_block:

Sequence 103, Application US/09358972

Patent No. 6235480

GENERAL INFORMATION:

APPLICANT: Shultz, John W

APPLICANT: Lewis, Martin K

APPLICANT: Lieppe, Donna

APPLICANT: Kephart, Michelle

APPLICANT: Rhodes, Richard B

APPLICANT: Olson, Ryan J

APPLICANT: Andriew, Christine A

APPLICANT: Hartnett, James R

APPLICANT: Gu, Trent

APPLICANT: Wood, Keith W

```

APPLICANT: Welch, Roy
TITLE OF INVENTION: Nucleic Acid Detection
FILE REFERENCE: PRO-103 6868/15528
CURRENT APPLICATION NUMBER: US/09/358,972
CURRENT FILING DATE: 1998-07-22
EARLIER APPLICATION NUMBER: 09/252,436
EARLIER FILING DATE: 1999-02-18
EARLIER APPLICATION NUMBER: 09/042,287
NUMBER OF SEQ ID NOS: 290
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO: 103
LENGTH: 30
TYPE: DNA
ORGANISM: Campylobacter jejuni
FEATURE:
OTHER INFORMATION: probe to Campylobacter jejuni
seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-406-147-32
seq_documentation_block:
sequence 34 Application US/09406147
; Sequence 34 Application US/09406147
; Patent No. 6270574
; GENERAL INFORMATION:
; APPLICANT: Shultz, John W
; APPLICANT: Lewis, Martin K
; APPLICANT: Manderkar, Michelle
; APPLICANT: Kephart, Daniel
; APPLICANT: Rhodes, Richard B
; APPLICANT: Andrews, Christine A
; APPLICANT: Hartnett, James R
; APPLICANT: Gu, Trent
; APPLICANT: Wood, Keith V
; APPLICANT: Welch, Roy
; TITLE OF INVENTION: EXOGENOUS NUCLEIC ACID DETECTION
; FILE REFERENCE: EXOGENOUS NUCLEIC ACID DETECTION
; CURRENT APPLICATION NUMBER: US/09/406,147
; CURRENT FILING DATE: 1998-09-27
; EARLIER APPLICATION NUMBER: 09/252,436
; EARLIER FILING DATE: 1999-02-18
; EARLIER APPLICATION NUMBER: 09/042,287
; EARLIER FILING DATE: 1998-03-13
; NUMBER OF SEQ ID NOS: 92
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO: 34
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Campylobacter jejuni
; US-09-406-147-34
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alignment_block:
US-09-439-311-2 x US-09-406-147-32/rev .. .
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97 GlnAspGlyGlnSerLeuLysThrArgThr 106
||||||||||||||||||||||||||||||||||| 1
1 CAAGATGGACAAAGTTAAACAGAACT 30
seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-09-130-663-22
seq_documentation_block:
sequence 22 Application US/09130663A
; Sequence 22 Application US/09130663A
; GENERAL INFORMATION:
; APPLICANT: Conklin, Darrell C.
; TITLE OF INVENTION: LIPOCALIN HOMOLOG
; FILE REFERENCE: 97-24
; CURRENT APPLICATION NUMBER: US/09/130,663A
; CURRENT FILING DATE: 1998-08-05
; EARLIER APPLICATION NUMBER: 60/054,867
; EARLIER FILING DATE: 1997-08-06
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO: 22
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alignment_block:
US-09-406-147-32
; Sequence 22 Application US/09130663A
; Patent No. 6020163
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Campylobacter jejuni
; US-09-406-147-32
seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-09-130-663-22
seq_documentation_block:
sequence 22 Application US/09130663A
; Sequence 22 Application US/09130663A
; GENERAL INFORMATION:
; APPLICANT: Conklin, Darrell C.
; TITLE OF INVENTION: LIPOCALIN HOMOLOG
; FILE REFERENCE: 97-24
; CURRENT APPLICATION NUMBER: US/09/130,663A
; CURRENT FILING DATE: 1998-08-05
; EARLIER APPLICATION NUMBER: 60/054,867
; EARLIER FILING DATE: 1997-08-06
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO: 22

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; LENGTH: 51
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION: Oligonucleotide primer: 2C13735.
; US-09-130-663-22

alignment_scores:
    Quality: 43.00      Length: 16
    Ratio: 3.583        Gaps: 0
    Percent Similarity: 75.000   Percent Identity: 50.000

alignment_block:
US-09-439-311-2 x US-09-130-663-22

Align seg 1/1 to: US-09-130-663-22 from: 1 to: 51

seq_name: /cgn2_6/ptodata/2/1na/6A_COMB.seq:US-09-081-180-29

seq_documentation_block:
Sequence 29, Application US/09081180
Patent No. 6022847

GENERAL INFORMATION:
APPLICANT: Sheppard, Paul O.

APPLICANT: Sheppard, Paul O.
; Sequence 29, Application US/09040786
; Patent No. 6025197

GENERAL INFORMATION:
APPLICANT: Sheppard, Paul O.

TITLE OF INVENTION: SECRETED SALIVARY ZSIG32
ADDRESS: ZymoGenetics
STREET: 1201 Eastlake Ave. E.
CITY: Seattle
STATE: WA
COUNTRY: USA
ZIP: 98102

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/081, 180
FILING DATE:
CLASSIFICATION:
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 60/041, 263
FILING DATE: March 19, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Lingenfelter, Susan E
REGISTRATION NUMBER: 41,156
REFERENCE/DOCKET NUMBER: 97-17
TELECOMMUNICATION INFORMATION:
TELEPHONE: 206-442-6675
TELEFAX: 206-442-6678

TELEX:
INFORMATION FOR SEQ ID NO: 29:
SEQUENCE CHARACTERISTICS:
LENGTH: 51 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
IMMEDIATE SOURCE:
CLONE: 2C13735
; INFORMATION FOR SEQ ID NO: 29:
SEQUENCE CHARACTERISTICS:
LENGTH: 51 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
IMMEDIATE SOURCE:
CLONE: 2C13735
; US-09-081-180-29

alignment_scores:
    Quality: 43.00      Length: 16
    Ratio: 3.583        Gaps: 0
    Percent Similarity: 75.000   Percent Identity: 50.000

alignment_block:
US-09-439-311-2 x US-09-081-180-29

Align seg 1/1 to: US-09-081-180-29 from: 1 to: 51

seq_name: /cgn2_6/ptodata/2/1na/6A_COMB.seq:US-09-040-786-29

seq_documentation_block:
Sequence 29, Application US/09040786
; Patent No. 6025197

GENERAL INFORMATION:
APPLICANT: Sheppard, Paul O.

TITLE OF INVENTION: SECRETED SALIVARY ZSIG32
ADDRESS: ZymoGenetics
STREET: 1201 Eastlake Ave. E.
CITY: Seattle
STATE: WA
COUNTRY: USA
ZIP: 98102

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/040, 786
FILING DATE:
CLASSIFICATION:
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 60/041, 263
FILING DATE: March 19, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Lingenfelter, Susan E
REGISTRATION NUMBER: 41,156
REFERENCE/DOCKET NUMBER: 97-17
TELECOMMUNICATION INFORMATION:
TELEPHONE: 206-442-6675
TELEFAX: 206-442-6678

TELEX:
INFORMATION FOR SEQ ID NO: 29:
SEQUENCE CHARACTERISTICS:
LENGTH: 51 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
IMMEDIATE SOURCE:
CLONE: 2C13735
; INFORMATION FOR SEQ ID NO: 29:
SEQUENCE CHARACTERISTICS:
LENGTH: 51 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
IMMEDIATE SOURCE:
CLONE: 2C13735
; US-09-081-180-29

Align seg 1/1 to: US-09-040-786-29 from: 1 to: 51

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seq\_name: /cgn2\_6/ptodata/2/1na/6A\_COMB.seq:US-09-432-335-22  
seq\_documentation\_block:  
Sequence 22, Application US/09432335  
GENERAL INFORMATION:  
PATENT NO.: 6143720  
APPLICANT: Conklin, Darrell C.  
TITLE OF INVENTION: LIPOCALIN HOMOLOG  
FILE REFERENCE: 97-24  
CURRENT APPLICATION NUMBER: US/09/432,335  
CURRENT FILING DATE: 1999-11-02  
EARLIER APPLICATION NUMBER: 09/130,663  
EARLIER FILING DATE: 1998-08-06  
EARLIER APPLICATION NUMBER: 60/054,867  
EARLIER FILING DATE: 1997-08-06  
NUMBER OF SEQ ID NOS: 30  
SOFTWARE: FASTSEQ for Windows Version 3.0  
SEQ ID NO: 22  
LENGTH: 51  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE: OTHER INFORMATION: Oligonucleotide primer: ZC13735  
us-09-432-335-22

alignment\_scores:  
Quality: 43.00 Length: 16  
Percent Similarity: 75.000 Percent Identity: 50.000

align seq 1/1 to: US-09-432-335-22 from: 1 to: 51

253 GlyValValIleGlyIysvaAspTywserAspGlyaspGluasnGly 268  
1 ||||| :||: ||||| :||| :||| :||| :||| :||| :||| :||| :||| :||| :|||  
seq\_name: /cgn2\_6/ptodata/2/1na/5A\_COMB.seq:US-08-219-012-80  
seq\_documentation\_block:  
Sequence 80, Application US/08219012  
Patent No. 5543293  
GENERAL INFORMATION:  
APPLICANT: Larry Gold  
APPLICANT: Diane Tasset  
TITLE OF INVENTION: Ligands of Thrombin  
NUMBER OF SEQUENCES: 92  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Swanson & Bratschun, L.L.C.  
STREET: 8400 E. Prentice Avenue, Suite 200  
CITY: Englewood  
STATE: Colorado  
COUNTRY: USA  
ZIP: 80111  
COMPUTER READABLE FORM:  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB storage  
COMPUTER: IBM compatible  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WordPerfect 6.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/687,421  
FILING DATE: 08-MAY-1996  
CLASSIFICATION: 435  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: 08/195,005  
FILING DATE: 10-FEBRUARY-1994  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: 07/973,333  
ATTORNEY/AGENT INFORMATION:

NAME: Barry J. Swanson  
REGISTRATION NUMBER: 33,215  
SEQUENCE/DOCKET NUMBER: 33,215  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (303) 850-9900  
TELEFAX: (303) 850-9401  
INFORMATION FOR SEQ ID NO: 80:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 60 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
US-08-219-012-80

alignment\_scores:  
Quality: 40.00 Length: 19  
Percent Similarity: 78.947 Percent Identity: 42.105

align seq 1/1 to: US-08-219-012-80 from: 1 to: 60

204 ThrSerValGlyIleGlyIleGlyAlaLeuAlaGluGluLeasnArgS 224  
1 |||:||:|||:|||:|||:|||:|||:|||:|||:|||:|||:|||:|||:|||:|||:|||:|||  
4 ACCGGGAGGGCGTAGGGTGGAGGCCATGGCTAGGCCACCGGA 53  
220 nalaASP 222  
:::||| 54 CTCGGAT 60

seq\_name: /cgn2\_6/ptodata/2/1na/6B\_COMB.seq:US-08-687-421-268  
seq\_documentation\_block:  
Sequence 268, Application US/08687421  
Patent No. 6177557  
GENERAL INFORMATION:  
APPLICANT: Gold, Larry  
APPLICANT: Janjic, Nebojsa  
APPLICANT: Tasset, Diane  
TITLE OF INVENTION: HIGH-AFFINITY LIGANDS OF BASIC  
TITLE OF INVENTION: FIBROBLAST GROWTH FACTOR AND  
TITLE OF INVENTION: THROMBIN  
NUMBER OF SEQUENCES: 445  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Swanson & Bratschun, L.L.C.  
STREET: 8400 E. Prentice Avenue, Suite 200  
CITY: Englewood  
STATE: Colorado  
COUNTRY: USA  
ZIP: 80111  
COMPUTER READABLE FORM:  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB storage  
COMPUTER: IBM compatible  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WordPerfect 6.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/687,421  
FILING DATE: 08-MAY-1996  
CLASSIFICATION: 435  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: 08/195,005  
FILING DATE: 10-FEBRUARY-1994  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: 07/973,333  
ATTORNEY/AGENT INFORMATION:



APPLICATION NUMBER: US 08/009, 266  
 FILING DATE: 22-JAN-1993  
 PRIORITY APPLICATION DATA:  
 APPLICATION NUMBER: US 07/894, 061  
 FILING DATE: 05-JUN-1992  
 PRIORITY APPLICATION DATA:  
 APPLICATION NUMBER: US 07/889, 724  
 FILING DATE: 26-MAY-1992  
 PRIORITY APPLICATION DATA:  
 ATTORNEY/AGENT INFORMATION:  
 NAME: Suh, Young J.  
 REGISTRATION NUMBER: P-41, 337  
 REFERENCE/DOCKET NUMBER: 27866/32760  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: (312) 474-6300  
 TELEFAX: (312) 474-0448  
 TELEX: (312) 474-6600  
 INFORMATION FOR SEQ ID NO: 97:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 47 base pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA  
 US-08-483-389-97

\* alignment\_scores:  
 Quality: 39.00 Length: 12  
 Ratio: 3.900 Gaps: 0  
 Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:  
 US-09-439-311-2 x US-08-483-389-97/rev . .

Align seq 1/1 to reverse of: US-08-483-389-97 from: 1 to: 47

169 ArgPheGluThrGlySerGlnSerPheSerSerGly 180  
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 44 AGGTGGAGACTGGTCAGCACGATTGGAGCGGA 9

seq\_name: /cgn2\_6/ptodata/2/1na/5B\_COMB.seq:US-08-487-113D-97

seq\_documentation\_block:  
 Sequence 97, Application US/08487113D  
 Patent No. 5837822

GENERAL INFORMATION:  
 APPLICANT: Gallatin, W. Michael  
 APPLICANT: Vazeux, Rosemary  
 TITLE OF INVENTION: ICAM-Related Materials and Methods  
 NUMBER OF SEQUENCES: 120

CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun  
 STREET: 6300 Sears Tower, 233 South Wacker Drive  
 CITY: Chicago  
 STATE: Illinois  
 COUNTRY: United States of America  
 ZIP: 60606-6402

COMPUTER READABLE FORM:  
 MEDIUM TYPE: floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/487,113D  
 FILING DATE: 05-AUG-1994  
 CLASSIFICATION: 424

PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: US 08/286, 754  
 FILING DATE: 07-JUN-1995  
 CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/102, 852  
 FILING DATE: 05-AUG-1993  
 PRIORITY APPLICATION DATA:  
 APPLICATION NUMBER: US 08/009, 266  
 FILING DATE: 22-JAN-1993  
 PRIORITY APPLICATION DATA:  
 APPLICATION NUMBER: US 07/894, 061  
 FILING DATE: 26-MAY-1992  
 PRIORITY APPLICATION DATA:  
 ATTORNEY/AGENT INFORMATION:  
 NAME: Gretta E.  
 REGISTRATION NUMBER: 35,302  
 REFERENCE/DOCKET NUMBER: 32744  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: (312) 474-6300  
 TELEFAX: (312) 474-0448  
 TELEX: 25-3856  
 INFORMATION FOR SEQ ID NO: 97:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 47 base pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA  
 US-08-487-113D-97

\* alignment\_scores:  
 Quality: 39.00 Length: 12  
 Ratio: 3.900 Gaps: 0  
 Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:  
 US-09-439-311-2 x US-08-487-113D-97/rev . .

Align seq 1/1 to reverse of: US-08-487-113D-97 from: 1 to: 47

169 ArgPheGluThrGlySerGlnSerPheSerSerGly 180  
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 44 AGGTGGAGACTGGTCAGCACGATTGGAGCGGA 9

seq\_name: /cgn2\_6/ptodata/2/1na/5B\_COMB.seq:US-08-473-503-97

seq\_documentation\_block:  
 Sequence 97, Application US/08473503  
 Patent No. 5892622

GENERAL INFORMATION:  
 APPLICANT: Gallatin, W. Michael  
 APPLICANT: Vazeux, Rosemary  
 TITLE OF INVENTION: ICAM-Related Materials and Methods  
 NUMBER OF SEQUENCES: 116

CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun  
 STREET: 6300 Sears Tower, 233 S. Wacker Drive  
 CITY: Chicago  
 STATE: Illinois  
 COUNTRY: USA  
 ZIP: 60606

COMPUTER READABLE FORM:  
 MEDIUM TYPE: floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/473,503  
 FILING DATE: 07-JUN-1995  
 CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/286,754  
 FILING DATE: 05-AUG-1994  
 APPLICATION NUMBER: US 08/102,852  
 FILING DATE: 05-AUG-1993  
 PRIORITY APPLICATION DATA:  
 APPLICATION NUMBER: US 08/009,266  
 FILING DATE: 22-JAN-1993  
 PRIORITY APPLICATION DATA:  
 APPLICATION NUMBER: US 07/894,051  
 FILING DATE: 05-JUN-1992  
 PRIORITY APPLICATION DATA:  
 APPLICATION NUMBER: US 07/889,724  
 FILING DATE: 26-MAY-1992  
 REFERENCE/DOCKET NUMBER: 33178  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: (312) 474-3300  
 TELEFAX: (312) 474-0448  
 TELEX: 25-3056  
 INFORMATION FOR SEQ ID NO: 97:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 47 base pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA  
 US-08-473-503-97  
  
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 Quality: 39.00      L  
 Percent Similarity: 83.333      Percent Ideepe:  
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 US-09-439-311-2 x US-08-473-503-97/rev  
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 seq\_documentation\_block:  
 Sequence 97, Application US/08483932  
 Patent No. 5880268  
 GENERAL INFORMATION:  
 APPLICANT: Galatin, W. Michael  
 APPLICANT: Vareux, Rosemary  
 TITLE OF INVENTION: ICAM Related Matri  
 NUMBER OF SEQUENCES: 116  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Marshall, O'Toole, Gers  
 STREET: 6300 Sears Tower, 233 S. W  
 CITY: Chicago  
 STATE: Illinois  
 COUNTRY: USA  
 ZIP: 60606  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: FLOPPY disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 SOFTWARE: PatentIn Release #1.0, V  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/483,932  
 FILING DATE: 07-JUN-1995

CLASSIFICATION: 530  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: 08/286,754  
FILING DATE: 05-AUG-1994  
APPLICATION NUMBER: US 08/102,852  
FILING DATE: 05-AUG-1993  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 08/009,266  
FILING DATE: 22-JAN-1993  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 07/894,061  
FILING DATE: 05-JUN-1992  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 07/889,724  
FILING DATE: 26-MAY-1992  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 07/827,689  
FILING DATE: 27-JAN-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: NO. 5880268band, Greta E.  
REGISTRATION NUMBER: 35,302  
REFERENCE/DOCKET NUMBER: 32178  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (312) 474-6300  
TELEFAX: (312) 474-0448  
TELEX: 25-3856  
INFORMATION FOR SEQ ID NO: 97:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA

US-08-183932-97

Wed Apr 17 07:36:49 2002

us-09-439-311-2.rni

Date: Apr 17, 2002 3:08 AM  
About: Results were produced by the Gencore software,  
Copyright (c) 1993-2000 Compugen Ltd.

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30	I	ARR3101	Sequence 103 from patient
50	I	A93165	Sequence 4 from Patient WC
51	I	A12223	01) nucleotide - 12/1993
51	I	A1224	01) nucleotide 12/1993
51	I	A1256	01) oligonucleotide 1/1994
51	I	A12597	oligonucleotide 1/1994
59	I	A01428	Sequence 105 from Patient
60	I	AR12592	Sequence 268 from patient
60	I	A12493	Sequence 80 from patient U
47	I	AR013897	Sequence 97 from patient
47	I	AR013851	Sequence 97 from patient
47	I	A0202511	Sequence 97 from patient
47	I	AR05831	Sequence 97 from patient
47	I	A0388217	Sequence 97 from patient
52	I	AF1033970	Saccharomyces cerevisiae
51	I	AX156673	Sequence 1 from Patent
51	I	AX156757	Sequence 2085 from Patent
51	I	AX158758	Sequence 2086 from Patent
51	I	AX158759	Sequence 2087 from Patent
53	I	AR0005607	Sequence 3 from patient
53	I	AR036067	Sequence 3 from patient
53	I	A125675	Sequence 3 from patient US
60	I	AX015223	Sequence 1 from patient
24	I	AR001228	Sequence 6 from patient
24	I	AR001521	Sequence 6 from patient
24	I	AR01078	Sequence 6 from patient
24	I	AR016102	Sequence 6 from patient
41	I	A18391	Sequence 6 from patient US
49	I	E13129	Process for producing now
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51	I	A1X58963	Sequence 2291 from Patent
51	I	A1X58964	Sequence 2292 from Patent
51	I	A1X61669	Sequence 4997 from Patent
53	I	A17409	Nucleotide sequence 15 fr
60	I	AR02802	Sequence 4 from patient
60	I	AR101743	Sequence 5 from patient
60	I	L139503	Homo sapiens (PB 3...PA635)
33	I	CA3210	Synthetic P1-P2 pdBP Link
34	I	AX131684	Sequence 17 from patient

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LOCUS	AR153100	30 bp	DNA	PAT	08-AUG-2001	
DEFINITION	Sequence 102 from patent US 62335480.					
ACCESSION	AR153100					
VERSION	AR153100.1					
KEYWORDS						
SOURCE	Unknown.					
ORGANISM	Unknown.					
UNCLASSIFIED	Unclassified.					
REFERENCE	1 (bases 1 to 30)					
AUTHORS	Shultz, J. William, Lewis, M. K., Leippe, D., Mandrekar, M., Kephart, D., Rhodes, R., Byron, Andrews, C. Ann, Hartnett, J. Robert, Gu, T., Olson, R. J., Wood, K. V. and Welch, R.					
TITLE	Detection of nucleic acid hybrids					
JOURNAL	Patent: US 6235480-A 102 22-MAY-2001;					
FEATURES	Location/Qualifiers					
source	1..30					
BASE COUNT	5 a 5 c	4 g	16 t			
ORIGIN						

US-09-439-311-2 x AR153101 ..

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97 GlnaspGivcInSerLeuLysThrArgThr 105  
 |||:|||||:|||||:|||||:|||||:  
 1 CAAGATGGCAAAAGTTAAAACAGAACT 30

seq\_name: gb\_pat:A93365

LOCUS	A93365	seq_documentation_block:	50 bp	DNA	PAT	22-JAN-2000
DEFINITION	Sequence 4 from Patent WO9744451.					
ACCESSION	A93365					
VERSION	A93365.1					
KEYWORDS	unidentified.					
ORGANISM	unclassified. 1 (bases 1 to 50)					
REFERENCE	Paesen,G.C. and Nuttall,P.A.					
AUTHORS						
TITLE	VASOACTIVE AMINE BINDING MOLECULES					
JOURNAL	PATENT: WO 974451-A 4 27-NOV-1997;					
FEATURES	OXFORD VACS LTD (GB); PAESEN GUIDO CHRISTIAN (GB) Location/Qualifiers					
SOURCE	1. .50 /organism="unidentified"					
BASE COUNT	11 a 8 c 14 g 17 t					
ORIGIN						

alignment\_scores:  
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 Percent Similarity: 100.000 Percent Identity: 70.000

alignment\_block:  
 US-09-439-311-2 x A93365 ..

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262 SeraspGlyAspGluAsnGlySerIleuIle 271  
 |||:|||||:|||||:|||||:  
 7 AGTGATGGATGATGATGGATCCCTCIG 36

seq\_name: gb\_pat:A12323

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DEFINITION	Oligonucleotide.					
ACCESSION	A12323					
VERSION	A12323.1					
KEYWORDS						
SOURCE	synthetic construct.					
ORGANISM	artificial sequence. 1 (bases 1 to 51)					
REFERENCE						
AUTHORS						
TITLE	HYBRID PROTEINS OR POLYPEPTIDES					
JOURNAL	PATENT: WO 8802157-A 24 21-APR-1988;					
FEATURES	Location/Qualifiers					
SOURCE	1. .51 /organism="synthetic construct"					
BASE COUNT	2 a 7 c 27 g 15 t					
ORIGIN						

alignment\_scores:  
 Quality: 41.00 Length: 17  
 Ratio: 3.154 Gaps: 0  
 Percent Similarity: 76.471 Percent Identity: 47.059

alignment\_block:  
 US-09-439-311-2 x A12324 ..

Align seg 1/1 to: A12324 from: 1 to: 51

300 GlyArgGlyIleLysIleThrGlySerIleGlyValGlyAlaGlyIle 316  
 |||:|||||:|||||:|||||:  
 1 GGGATGGGGTTGGCGTGGGGTGGCGTTGGGGTGGCGTTGGATCCT 50  
 316 u 316  
 51 c 51

seq\_name: gb\_pat:A12596

LOCUS	A12596	seq_documentation_block:	51 bp	DNA	PAT	05-JAN-1994
DEFINITION	Oligonucleotide.					
ACCESSION	A12596					
VERSION	A12596.1					
KEYWORDS						
SOURCE	synthetic construct.					
ORGANISM	synthetic construct. artificial sequence. 1 (bases 1 to 51)					
REFERENCE						
AUTHORS						
TITLE	RECOMBINANT VIRUS					
JOURNAL	PATENT: WO 8701386-A 12 12-MAR-1987;					
FEATURES	Location/Qualifiers					
SOURCE	1. .51 /organism="synthetic construct"					
BASE COUNT	15 a 27 c 7 g 2 t					
ORIGIN						

alignment\_scores:  
 Quality: 41.00 Length: 17  
 Ratio: 3.154 Gaps: 0  
 Percent Similarity: 76.471 Percent Identity: 47.059

BASE COUNT 15 a 27 c 7 g 2 t ORIGIN /db\_xref="taxon:32630"

alignment\_scores: Quality: 41.00 Length: 17 Gaps: 0 Percent Similarity: 76.471 Percent Identity: 47.059

alignment\_block: US-09-439-311-2 x A12596/rev

Align seg 1/1 to reverse of: A12596 from: 1 to: 51

300 GlyArgGlyIleLysIleThrGlySerIleGlyValGlyAlaGlyIleLe 316  
 51 GGGATCGGGTGTGGCTTGCGTTGGGAGTCCT 2

316 u 316  
 1 C 1

seq\_name: gb\_pat:A12597

seq\_documentation\_block:  
 LOCUS A12597 51 bp DNA PAT 05-JAN-1994  
 DEFINITION Oligonucleotide.  
 ACCESSION A12597  
 VERSION A12597.1 GI:489543  
 KEYWORDS synthetic construct.  
 SOURCE synthetic construct.  
 ORGANISM synthetic construct.  
 FEATURES artificial sequence.  
 REFERENCE 1 (bases 1 to 51)  
 AUTHORS RECOMBINANT VIRUS  
 JOURNAL Patent: WO 9701386 A 13 12-MAR-1987;  
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 /db\_xref="taxon:32630"

BASE COUNT 2 a 7 c 27 g 15 t ORIGIN

alignment\_scores: Quality: 40.00 Length: 15 Gaps: 0 Percent Similarity: 86.667 Percent Identity: 53.333

alignment\_block: US-09-439-311-2 x AX011428

Align seg 1/1 to: AX011428 from: 1 to: 59

163 SerLysIleGlyValThrArgPheGluThrGlySerGlnSerPhe 177  
 9 AGCTCTTGGGTTGGCGGTTTCCACTGGCAGCTGAAGTC 53

seq\_name: gb\_pat:AR125926

seq\_documentation\_block:  
 LOCUS AR125926 60 bp DNA PAT 16-MAY-2001  
 DEFINITION Sequence 268 from Patent US 6177557.  
 ACCESSION AR125926  
 ACCESSION AR125926.1 GI:14111988  
 VERSION AR125926.1  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unclassified.  
 FEATURES  
 REFERENCE 1 (bases 1 to 60)  
 AUTHORS TITLE Janjic,N., Gold,L. and Tasset,D.  
 JOURNAL High affinity ligands of basic fibroblast growth factor and  
 FEATURES source thrombin Patent: US 6177557-A 268 23-JAN-2001;  
 BASE COUNT 10 a 11 c 29 g 10 t ORIGIN

align seg 1/1 to: A12597 from: 1 to: 51

300 GlyArgGlyIleLysIleThrGlySerIleGlyValGlyAlaGlyIleLe 316  
 1 GGCATCGGGTGTGGCTTGCGTTGGGAGTCCT 50

316 u 316  
 1 C 51

seq\_name: gb\_pat:AX011428

seq\_documentation\_block:  
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 DEFINITION Sequence 105 from Patent WO9955907.  
 ACCESSION AX011428  
 VERSION AX011428.1 GI:9997978  
 KEYWORDS synthetic construct.

Align seg 1/1 to: AR125926 from: 1 to: 60

204 ThrSerValGlyThrGlyLeuGlyAlaLeuAlaGluGluLeAsnArgAs 220  
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 220 nalaAsp 222  
 54 CTCGGAT 60

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seq_name: gb_pat:I24293
seq_documentation_block: I24293 60 bp DNA
DEFINITION Sequence 80 from patent US 5543293.
ACCESSION I24293
Locus KEYWORDS Unknown.
ORGANISM Unclassified.
REFERENCE 1. (bases 1 to 60)
AUTHORS Gold,L. and Tasset,D.
TITLE DNA ligands of thrombin
JOURNAL Patent: US 5543293 A 80 06-AUG-1996;
FEATURES source 1. . 60
BASE COUNT 10 a 10 t
ORIGIN 10 a /organism="unknown"
11 c 29 g 10 t

alignment_scores: Quality: 40.00 Length: 19
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Percent Identity: 78.947 Percent Identity: 42.105

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US-09-439-311-2 x I24293 . .

alignment_block: 204 ThrSerValGlyThrGlyLeuGlyAlaLeuAlaGluGlutLeasnArgAs 220
4 ACCGGGGAGGCGTAGGGTGGAGGCCGATGTGGTAGGCACCGA 53
220 nalaAsp 222
54 CTCGGAT 60

seq_name: gb_pat:AR013897

seq_documentation_block: AR013897 47 bp DNA
DEFINITION Sequence 97 from patent US 5773218.
ACCESSION AR013897
VERSION AR013897.1 GI:3971351
KEYWORDS SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1. (bases 1 to 47)
AUTHORS Gilliam,W.Michael and Vazeux,R.
TITLE Method to identify compounds which modulate ICAM-related protein
interactions
JOURNAL Patent: US 5773218 A 97 30-JUN-1998;
FEATURES source 1. . 47
BASE COUNT 9 a 10 t
ORIGIN 9 a /organism="unknown"
21 c 7 g 10 t

alignment_scores: Quality: 39.00 Length: 12
Percent Similarity: 3.900 Gaps: 0
Percent Identity: 83.333 Percent Identity: 66.667

alignment_block: US-09-439-311-2 x AR013897/rev . .

Align seg 1/1 to reverse of: AR013897 from: 1 to: 47

seq_name: gb_Pat:AR042511
seq_documentation_block: AR042511 47 bp DNA
DEFINITION Sequence 97 from patent US 5811517.
ACCESSION AR042511
VERSION AR042511.1 GI:5963007
KEYWORDS SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1. (bases 1 to 47)
AUTHORS Gilliam,W.Michael and Vazeux,R.
TITLE ICAM-related protein variants
JOURNAL Patent: US 5811517 A 97 22-SEP-1998;
FEATURES source 1. . 47
BASE COUNT 9 a 10 t
ORIGIN 9 a /organism="unknown"
21 c 7 g 10 t

alignment_scores: Quality: 39.00 Length: 12
Percent Similarity: 3.900 Gaps: 0
Percent Identity: 83.333 Percent Identity: 66.667

alignment_block: US-09-439-311-2 x AR042511/rev . .

Align seg 1/1 to reverse of: AR042511 from: 1 to: 47

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seq\_name: gb\_pat:AR058391

seq\_documentation\_block:

LOCUS AR058391 47 bp DNA

DEFINITION Sequence 97 from patent US 5837822.

ACCESSION AR058391

VERSION AR058391.1 GI:5983968

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 47)

AUTHORS Gallatin,W Michael and Vazeux,R.

JOURNAL Patent: US 5837822-A 97-17-NOV-1998;

FEATURES Location/Qualifiers

source

BASE COUNT 9 a /organism="unknown"

ORIGIN 21 c 7 g 10 t

alignment\_scores:

Quality: 39.00 Length: 12

Ratio: 3.900 Gaps: 0

Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:

US-09-439-311-2 x AR058391/rev

Align seg 1/1 to reverse of: AR058391 from: 1 to: 47

169 ArgPheGluThrGlySerGlnSerPheSerSerGly 180  
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 44 AGGATGGAGACTGGTCAGCACCATTTGGAGCTGGA 9

seq\_name: gb\_pat:AR088217

seq\_documentation\_block:

LOCUS AR088217 47 bp DNA

DEFINITION Sequence 97 from patent US 5989843.

ACCESSION AR088217

VERSION AR088217.1 GI:10014980

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 47)

AUTHORS Gallatin,W Michael and Vazeux,R.

TITLE Methods for identifying modulators of protein kinase C phosphorylation of ICAM-related protein

Patent: US 5989843-A 97-23-NOV-1999;

FEATURES Location/Qualifiers

source 1..47

BASE COUNT 9 a /organism="unknown"

ORIGIN 21 c 7 g 10 t

alignment\_scores:

Quality: 39.00 Length: 12

Ratio: 3.900 Gaps: 0

Percent Similarity: 83.333 Percent Identity: 66.667

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seq\_name: gb\_pat:AR088217

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LOCUS AR088217 47 bp DNA

DEFINITION Sequence 97 from patent US 5989843.

ACCESSION AR088217

VERSION AR088217.1 GI:10014980

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 47)

AUTHORS Gallatin,W Michael and Vazeux,R.

JOURNAL Patent: US 5989843-A 97-23-NOV-1999;

FEATURES Location/Qualifiers

source 1..47

BASE COUNT 9 a /organism="unknown"

ORIGIN 21 c 7 g 10 t

alignment\_scores:

Quality: 39.00 Length: 12

Ratio: 3.900 Gaps: 0

Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:

US-09-439-311-2 x AR088217/rev

Align seg 1/1 to reverse of: AR088217 from: 1 to: 47

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seq\_name: gb\_pat:AR088217

seq\_documentation\_block:

LOCUS AR088217 47 bp DNA

DEFINITION Sequence 97 from patent US 5989843.

ACCESSION AR088217

VERSION AR088217.1 GI:10014980

KEYWORDS

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 47)

AUTHORS Gallatin,W Michael and Vazeux,R.

JOURNAL Patent: US 5989843-A 97-23-NOV-1999;

FEATURES Location/Qualifiers

source 1..47

BASE COUNT 9 a /organism="unknown"

ORIGIN 21 c 7 g 10 t

Wed Apr 17 07:36:47 2002

us-09-439-311-2.rge

Gencore version 4.5  
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On nucleic - nucleic search, using sw model  
Run on: April 16, 2002, 23:26:24 ; Search time 1531.8 Seconds  
(without alignments)  
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Title: US-09-439-311-1

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Searched: 1472140 seqs, 8248589755 residues

Total number of hits satisfying chosen parameters: 586436

Minimum DB seq length: 0

Maximum DB seq length: 60

Post-processing: Minimum Match 0%

Listing first 45 summaries

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c 4	20.8	2.1	24	6	AR001228	AR001228 Sequence
c 5	20.8	2.1	24	6	AR00251	AR00251 Sequence
c 6	20.8	2.1	24	6	AR010178	AR010178 Sequence
c 7	20.8	2.1	24	6	AR061102	AR061102 Sequence
c 8	20.8	2.1	24	6	I38291	I38291 Sequence 6
c 9	20.8	2.1	44	6	AX052949	AX052949 Sequence 6
c 10	20.8	2.1	58	6	AR061278	AR061278 Sequence
c 11	20.6	2.1	60	6	AB0405	AB0405 Sequence 17
c 12	20.4	2.0	51	6	AX160493	AX160493 Sequence
c 13	20.4	2.0	51	6	AX160494	AX160494 Sequence
c 14	20.4	2.0	57	3	AF320169	AF320169 Drosophil
c 15	20.4	2.0	57	3	AF320170	AF320170 Drosophil
c 16	20.4	2.0	57	3	AF320171	AF320171 Drosophil
c 17	20.4	2.0	57	3	AF320172	AF320172 Drosophil
c 18	20.4	2.0	57	3	AF320173	AF320173 Drosophil
c 19	20.4	2.0	57	3	AF320174	AF320174 Drosophil
c 20	20.4	2.0	57	3	AF320175	AF320175 Drosophil
c 21	20.4	2.0	57	3	AF320176	AF320176 Drosophil
c 22	20.4	2.0	57	3	AF320177	AF320177 Drosophil
c 23	20.4	2.0	57	3	AF320178	AF320178 Drosophil
c 24	20.4	2.0	57	3	AF320179	AF320179 Drosophil
c 25	20.2	2.0	52	6	A69988	A69988 Sequence 19
c 26	20.2	2.0	50	6	A17107	A17107 Oligonucleo
c 27	20.2	2.0	50	6	AR027492	AR027492 Sequence
c 28	20.2	2.0	57	6	AR07133	AR07133 Sequence
c 29	20.2	2.0	57	6	AR102800	AR102800 Sequence
c 30	20.2	2.0	57	6	I21136	I21136 Sequence 4
c 31	20.2	2.0	60	6	A38623	A38623 Sequence 16
c 32	20.2	2.0	60	6	AR040717	AR040717 Sequence
c 33	19.8	2.0	51	6	AX165291	AX165291 Sequence
c 34	19.8	2.0	59	6	AX011435	AX011435 Sequence
c 35	19.6	2.0	29	6	A20557	A20557 Oligonucleo
c 36	19.6	2.0	43	6	I11627	I11627 Sequence 12
c 37	19.6	2.0	51	6	AX160235	AX160235 Sequence
c 38	19.6	2.0	55	14	HIV045041	U45041 Human Immuno
c 39	19.6	2.0	57	6	A32988	A32988 Synthetic P
c 40	19.6	2.0	58	6	AR011305	AR011305 Sequence
c 41	19.6	2.0	58	6	I17943	I17943 Sequence 17
c 42	19.4	1.9	37	6	AR06304	AR06304 Sequence
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c 44	19.4	1.9	51	6	AX117493	AX117493 Sequence
c 45	19.4	1.9	51	6	AX162140	AX162140 Sequence

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RESULT	1	AR153100/C	LOCUS	AR153100	30 bp	DNA	PAT	08-AUG-2001
DEFINITION			SEQUENCE	Sequence 102 from patent US 6235480.				
ACCESSION			DEFINITION	AR153100				
VERSION			ACCESSION	AR153100.1	GI:15120632			
KEYWORDS			KEYWORDS					
SOURCE	Unknown.		ORGANISM	Unclassified.				
REFERENCE	1 (bases 1 to 30)		REFERENCE	Shultz,J.William, Lewis,M.K., Leippe,D., Mandrekar,M., Rephart,D., Olson,R.J., Wood,K.V. and Welch,R.				
AUTHORS	Rhodes,R.Byron, Andrews,C.Ann, Hartnett,J.Robert, Gu,T.,		JOURNAL	Detection of nucleic acid hybrids				
FEATURES	Patent: US 6235480-A 102-22-MAY-2001;		FEATURES	Location/Qualifiers				
SOURCE	1. 30		SOURCE	/organism="unknown"				
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PAT

08-AUG-2001

BASE COUNT

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7 t

1 others

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DB 0; Mismatches

0; Indels

0; Gaps

0; Matches

0; Accession

AR153101.1

GI:15120633

KEYWORDS

ORGANISM

Unknown.

SOURCE

Unclassified.

REFERENCE

1 (bases 1 to 30)

AUTHORS

Rhodes, R., Bryon, Andrews, C., Ann, Hartnett, J., Robert, Gu, T.,

Olson, R.J., Wood, K.V., and Welch, R.

TITLE

Detection of nucleic acid hybrids

JOURNAL

Patent: US 6235480-A 103 22-MAY-2001;

FEATURES

Location/Qualifiers

1. .30

/organism="unknown"

ORIGIN

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Query Match

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AR153101

GI:15120633

KEYWORDS

ORGANISM

Unknown.

Query Match

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Length 30;

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0; Gaps

0; Matches

0; Accession

AR153101

GI:15120633

KEYWORDS

ORGANISM

Unknown.

Query Match

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Length 30;

DB 0; Mismatches

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AR153101

GI:15120633

KEYWORDS

ORGANISM

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Query Match

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KEYWORDS

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Query Match

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KEYWORDS

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Query Match

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Query Match

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0; Matches

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AR153101

GI:15120633

KEYWORDS

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Query Match

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KEYWORDS

ORGANISM

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Query Match

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Length 30;

DB 0; Mismatches

FEATURES	JOURNAL	Patent: US 5753444-A 6 19-MAY-1998;
source	Location/Qualifiers	
BASE COUNT	6 a 5 c 5 g 8 t	
ORIGIN		
RESULT	8	
QY	91 ggctttagaatcaactccggacca 114	
LOCUS	AR010178/c	
DEFINITION	Best Local Similarity 91.7%; Pred. No. 1.9e+06; Sequence 6 from patent US 5756701.	
ACCESSION	I38291	
VERSION	I38291.1	
KEYWORDS	GI:3968983	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE	1 (bases 1 to 24)	
AUTHORS	Wu, L., Coombs,J., Malmstrom,S.L. and Glass,M.J.	
TITLE	Specific oligonucleotide primer pairs and probes for discriminating	
FEATURES	specific analytes	
JOURNAL	Patent: US 5756701-A 6 26-MAY-1998;	
source	Location/Qualifiers	
BASE COUNT	6 a 5 c 5 g 8 t	
ORIGIN		
RESULT	9	
QY	91 ggctttagaatcaactccggacca 114	
LOCUS	AR010178/c	
DEFINITION	Best Local Similarity 91.7%; Pred. No. 1.9e+06; Sequence 6 from patent US 5756701.	
ACCESSION	I38291	
VERSION	I38291.1	
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SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE	1 (bases 1 to 24)	
AUTHORS	Wu, L., Coombs,J., Malmstrom,S.L. and Glass,M.J.	
TITLE	Specific oligonucleotide primer pairs and probes for discriminating	
FEATURES	specific analytes	
JOURNAL	Patent: US 5756701-A 6 26-MAY-1998;	
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BASE COUNT	6 a 5 c 5 g 8 t	
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VERSION	I38291.1	
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ORGANISM	Unclassified.	
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AUTHORS	Wu, L., Coombs,J., Malmstrom,S.L. and Glass,M.J.	
TITLE	Specific oligonucleotide primer pairs and probes for discriminating	
FEATURES	specific analytes	
JOURNAL	Patent: US 5756701-A 6 26-MAY-1998;	
source	Location/Qualifiers	
BASE COUNT	6 a 5 c 5 g 8 t	
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LOCUS	AR010178/c	
DEFINITION	Best Local Similarity 91.7%; Pred. No. 1.9e+06; Sequence 6 from patent US 5756701.	
ACCESSION	I38291	
VERSION	I38291.1	
KEYWORDS	GI:2086281	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE	1 (bases 1 to 24)	
AUTHORS	Wu, L., Coombs,J., Malmstrom,S.L. and Glass,M.J.	
TITLE	Methods, kits and solutions for preparing sample material for	
FEATURES	nucleic acid amplification	
JOURNAL	Patent: US 5612473-A 6 18-MAR-1997;	
source	Location/Qualifiers	
BASE COUNT	6 a 5 c 5 g 8 t	
ORIGIN		
RESULT	9	
QY	91 ggctttagaatcaactccggacca 114	
LOCUS	AR010178/c	
DEFINITION	Best Local Similarity 91.7%; Pred. No. 1.9e+06; Sequence 6 from patent US 5756701.	
ACCESSION	I38291	
VERSION	I38291.1	
KEYWORDS	GI:12227051	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE	1 (bases 1 to 44)	
AUTHORS	Kwagh,J.G., Macklin,J.J., Mitsis,P.G. and Ulmer,K.M.	
TITLE	Methods for sequencing and characterizing polymeric biomolecules using	
FEATURES	aptamers and a method for producing aptamers	
JOURNAL	Patent: WO 0071755-A 55 30-NOV-2000;	
source	Praelux Incorporated (US)	
BASE COUNT	9 a 6 c 19 g 10 t	
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QY	747 tgcatcaatgggttgataaqtgaatgtttatcca 786	
LOCUS	AR010178/c	
DEFINITION	Best Local Similarity 91.7%; Pred. No. 1.9e+06; Sequence 6 from patent US 5756701.	
ACCESSION	I38291	
VERSION	I38291.1	
KEYWORDS	GI:2086281	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE	1 (bases 1 to 24)	
AUTHORS	Wu, L., Coombs,J., Malmstrom,S.L. and Glass,M.J.	
TITLE	Methods and apparatus for preparing, amplifying, and discriminating	
FEATURES	multiple analytes	
JOURNAL	Patent: US 5846783-A 6 08-DEC-1998;	
source	Location/Qualifiers	
BASE COUNT	6 a 5 c 5 g 8 t	
ORIGIN		
RESULT	10	
QY	747 tgcatcaatgggttgataaqtgaatgtttatcca 786	
LOCUS	AR010178/c	
DEFINITION	Best Local Similarity 91.7%; Pred. No. 1.9e+06; Sequence 6 from patent US 5756701.	
ACCESSION	I38291	
VERSION	I38291.1	
KEYWORDS	GI:2086281	
SOURCE	Unknown.	
ORGANISM	Unclassified.	
REFERENCE	1 (bases 1 to 24)	
AUTHORS	Wu, L., Coombs,J., Malmstrom,S.L. and Glass,M.J.	
TITLE	Methods and apparatus for preparing, amplifying, and discriminating	
FEATURES	multiple analytes	
JOURNAL	Patent: US 5846783-A 6 08-DEC-1998;	
source	Location/Qualifiers	
BASE COUNT	6 a 5 c 5 g 8 t	
ORIGIN		

LOCUS	DEFINITION	SEQUENCE	PAT	29-SEP-1999
AR061278	Sequence 7 from Patent US 5843650.			
ACCESSION	AR061278		AUTHORS	Shimmetts,R.A. and Reach,M.
VERSION	AR061278.1		TITLE	Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
KEYWORDS			JOURNAL	Patent: WO 0140521-A 3821 07-JUN-2001;
ORGANISM	Unknown.		CURATOR	Coragen Corporation (US)
REFERENCE	Unclassified.		FEATURES	Location/Qualifiers
AUTHORS	1 (bases 1 to 58)		source	1. .51 /organism="Homo sapiens" /ab_xref="taxon:9606"
TITLE	Segev,D.			/note="1 of 2 allelic variants (3821 is other entry) Accession number cg43921050"
JOURNAL	Nucleic acid detection and amplification by chemical linkage of			
FEATURES	Patent: US 5843650-A 7 01-DEC-1998;			
SOURCE	Location/Qualifiers			
BASE COUNT	15 a 18 c 13 g 12 t			
ORIGIN				
RESULT	11			
LOCUS	A80405	60 bp DNA	PAT	21-JAN-2000
DEFINITION	Sequence 17 from Patent WO951771.			
ACCESSION	A80405			
VERSION	1 GI:6731293			
KEYWORDS				
ORGANISM	Archaeoglobus fulgidus.			
REFERENCE	Archaeoglobus fulgidus			
AUTHORS	Euryarchaeota; Archaeoglobales; Archaeoglobaceae;			
TITLE	Archaeoglobus			
JOURNAL	1 (bases 1 to 60)			
FEATURES	Jansen, R. and Schouts, L.M.			
SOURCE	A METHOD OF INVERSTRAIN DIFFERENTIATION OF BACTERIA			
BASE COUNT	20 a 11 c 9 g 20 t			
ORIGIN				
RESULT	12			
LOCUS	AX160493	51 bp DNA	PAT	22-JUN-2001
DEFINITION	Sequence 3821 from Patent WO0140521.			
ACCESSION	AX160493			
VERSION	AX160493.1 GI:14541824			
KEYWORDS	human.			
SOURCE	Homo sapiens			
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; vertebrata; Euteleostomi; Mammalia; Buterilia; Primates; Catarrhini; Hominidae; Homo.			
RESULT	13			
LOCUS	AX160494	51 bp DNA	PAT	22-JUN-2001
DEFINITION	Sequence 3822 from Patent WO0140521.			
ACCESSION	AX160494			
VERSION	AX160494.1 GI:14541825			
KEYWORDS	human.			
ORGANISM	Homo sapiens			
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; vertebrata; Euteleostomi; Mammalia; Buterilia; Primates; Catarrhini; Hominidae; Homo.			
AUTHORS	1 (bases 1 to 51)			
TITLE	Shimmetts, R.A. and Leach, M.			
JOURNAL	Nucleic acids containing single nucleotide polymorphisms and methods of use thereof			
FEATURES	Patent: WO 0140521-A 3822 07-JUN-2001;			
SOURCE	Curagen Corporation (US)			
BASE COUNT	20 a 18 c 5 g 20 t			
ORIGIN				
RESULT	14			
LOCUS	AF320169	57 bp DNA	INV	23-APR-2001
DEFINITION	Drosophila pseudoobscura strain Mather10 bicoid (bcd) gene, partial cds.			
ACCESSION	AF320169			
VERSION	AF320169.1 GI:13752326			
KEYWORDS				
SOURCE	Drosophila pseudoobscura.			
ORGANISM	Drosophila pseudoobscura.			



Wed Apr 17 07:36:44 2002

us-09-439-311-1.rge

GenCore version 4.5									
Copyright (c) 1993 - 2000 Compugen Ltd.									
OM nucleic - nucleic search, using sw model									
Run on:	April 17, 2002, 01:18:44 ; Search time 172.86 Seconds (without alignments)								
Title:	US-09-439-311-1								
Perfect score:	999								
Sequence:	1 attaacacaatgttcagc.....taaaaaatgtatgttagat 999								
Scoring table:	IDENTITY_NUC								
Gapop 10.0 , Gapext 1.0									
Searched:	930621 seqs, 428662619 residues								
Total number of hits satisfying chosen parameters:	1026190								
Minimum DB seq length: 0									
Maximum DB seq length: 60									
Post-processing: Minimum Match 0%									
Listing first 45 summaries									
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.									
SUMMARIES									
Result NO.	Score	Query Length	DB ID	Description					
C 1	25.8	2.7	30 21 AAB8691	Probe to Campylobacter jejuni	XX	PR	18-FEB-1999; 990US-0252436.	AAV31446	Campylobacter nucl
C 2	26.8	2.7	30 21 AA86892	Probe to Campylobacter	XX	PR	21-JUL-1999; 990US-0358972.	AAV5942	Oligonucleotide PC
C 3	26.8	2.7	30 21 AA93189	Campylobacter jejuni	XX	PR	25-AUG-1999; 990US-0383316.	AAV20847	Campylobacter CRO4
C 4	26.8	2.7	30 21 AA93190	Campylobacter jejuni	XX	PR	44-22 AAF6711	deKb-Specific apta	
C 5	21.8	2.2	50 20 AA52169	Synthetic plasmid	XX	PA	(PROM-) PROMEGA CORP.	AAV69291	Staphylococcus aur
C 6	21.4	2.1	51 19 AAV04223	Human cardiac trop	XX	PI	Shultz JW, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;	AAV15200	
C 7	21.4	2.1	58 19 AAV04244	Human cardiac trop	XX	PI	Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;	AAV207004	
C 8	21.2	2.1	54 21 AA97418	beta wild-type pRZ	XX	PT		AAV16354	
C 9	21	2.1	27 21 AA27148	Campylobacter coli	XX	DR	WPI: 2000-56537752.	AAV160503	
C 10	21	2.1	33 21 AA27149	Campylobacter coli	XX	PR			
C 11	20.8	2.1	24 18 AAV160503	Primer CFO4R.4.	XX	PR			

PT Determining presence or absence of a predetermined endogenous nucleic acid sequence by using an enzyme that depolymerizes the 3' end of an oligonucleotide probe hybridized to a target sequence to release identifier nucleotides -

XX Example; Page 321; 389pp; English.

CC The present invention describes a method (M1) for determining the presence or absence of a predetermined endogenous nucleic acid target sequence (ENAT). The method comprises hybridising a probe having an identifier nucleotide (IN) with ENAT which is treated with an enzyme that depolymerises the 3' end of hybridised NA to release the INs.

CC M1 is used for determining the number of known sequence repeats present in a nucleic acid target sequence in a nucleic acid sample. The method is also useful for determining whether a nucleic acid target sequence in a sample is an allele from a homozygous or heterozygous locus. The method is also useful for detection of mutations, translocations and SNPs in nucleic acids (including those associated with genetic disease), contamination, and analysis of forensic samples. AA86791 to AAB12817 represent sequence which are used in the exemplification of the present invention.

N.B. There is a discrepancy between the SEQ ID NO: and sequences given in the examples, and the SEQ ID NO: and sequences given in the sequence listing from the present invention.

CC Sequence 30 BP; 5 A; 5 C; 4 G; 16 T; 0 other;

XX Query Match 2.7%; Score 26.8; DB 21; Length 30; Best Local Similarity 93.3%; Pred. No. 1.3e+03; Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 289 caagatgttcaaaacttaaaaaaagaact 318

Db 30 CAAGATGGACAAGTTAAAAACAAACT 1

RESULT 2

ID AA86892

ID AA86892 standard; DNA; 30 BP.

XX AC AA86892;

XX DT 15-JAN-2001 (first entry)

XX DE Probe to *Campylobacter jejuni*.

XX KW Detection; nucleic acid hybrid; depolymerisation; analysis; SNP; single nucleotide polymorphism; identification; viral load; probe; genotyping; medical marker diagnostic; primer; target; mutation; genetic disease; ss.

XX OS *Campylobacter jejuni*.

XX PN WO200049180-A1.

XX PD 24-AUG-2000.

XX PF 1B-FEB-2000; 2000WO-US04242.

XX PR 1B-FEB-1999; 99US-0252436.

XX PR 21-JUL-1999; 99US-0358972.

XX PR 25-AUG-1999; 99US-0383316.

XX PA (PROM-) PROMEGA CORP.

XX PI Shultz JW, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB; Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;

XX PT WPI; 2000-565377/52.

PT Determining presence or absence of a predetermined endogenous nucleic

acid sequence by using an enzyme that depolymerizes the 3' end of an oligonucleotide probe hybridized to a target sequence to release identifier nucleotides -

XX Example; Page 321; 389pp; English.

CC The present invention describes a method (M1) for determining the presence or absence of a predetermined endogenous nucleic acid target sequence (ENAT). The method comprises hybridising a probe having an identifier nucleotide (IN) with ENAT which is treated with an enzyme that depolymerises the 3' end of hybridised NA to release the INs.

CC M1 is used for determining the number of known sequence repeats present in a nucleic acid target sequence in a nucleic acid sample. The method is also useful for determining whether a nucleic acid target sequence in a sample is an allele from a homozygous or heterozygous locus. The method is also useful for detection of mutations, translocations and SNPs in nucleic acids (including those associated with genetic disease), determination of viral load, species identification, sample contamination, and analysis of forensic samples. AA86791 to AAB12817 represent sequence which are used in the exemplification of the present invention.

N.B. There is a discrepancy between the SEQ ID NO: and sequences given in the examples, and the SEQ ID NO: and sequences given in the sequence listing from the present invention.

XX Sequence 30 BP; 16 A; 4 C; 5 G; 5 T; 0 other;

XX Query Match 2.7%; Score 26.8; DB 21; Length 30; Best Local Similarity 93.3%; Pred. No. 1.3e+03; Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 289 caagatgttcaaaacttaaaaaaagaact 318

Db 1 caagatggacaagttaaaaacaagaact 30

RESULT 3

ID AA93188/C

ID AA93188 standard; DNA; 30 BP.

XX AC AAA93188;

XX DT 11-JAN-2001 (first entry)

XX DE *Campylobacter jejuni* interrogation probe 11451.

XX KW *Campylobacter jejuni*; nucleic acid detection; genomic typing; mutation detection; viral load determination; species identification; forensic analysis; probe; ss.

XX OS *Campylobacter jejuni*.

XX PN WO200049179-A1.

XX PD 24-AUG-2000.

XX PF 1B-FEB-2000; 2000WO-US04176.

XX PR 1B-FEB-1999; 99US-0252436.

XX PR 21-JUL-1999; 99US-0358972.

XX PR 27-SEP-1999; 99US-0406147.

XX PA (PROM-) PROMEGA CORP.

XX PI Shultz JW, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB; Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;

XX DR WPI; 2000-549282/50.

PT Detecting the presence of predetermined exogenous nucleic acid target sequence useful for e.g. genotyping, comprises depolymerizing the 3' end of an oligonucleotide probe hybridized to a nucleic acid target

PT sequence " XX  
 PS Claim 47; Page 187; 230pp; English.  
 XX  
 CC The present sequence is an interrogation probe which was used to detect a  
 CC segment of the genome of *Campylobacter jejuni*. This was performed as part  
 CC of a method for determining the presence of a known exogenous nucleic  
 CC acid target sequence in a nucleic acid sample. The method comprises  
 CC adding a treated sample with a depolymerising enzyme which releases one  
 CC or more nucleotides from the 3'-end of a hybridised nucleic acid probe.  
 CC The method is used for assaying nucleic acids for a particular native or  
 CC mutant sequence, and for genomic typing. It is useful for detecting  
 CC mutations, translocations, and single nucleotide polymorphisms,  
 CC determination of viral load, species identification, detection of sample  
 CC contamination, and analysis of forensic samples. Compared with previous  
 CC methods of detecting nucleic acid hybrids, the new method has higher  
 CC sensitivity without the need for radiochemicals or electrophoresis. It is  
 CC quantitatively, highly reproducible and can be automated. The method can  
 CC reliably detect as few as 10 copies of a virus in a sample, and is  
 CC capable of providing multiple analyses in a single assay (multiplex  
 XX assay).  
 Sequence 30 BP; 5 A; 5 C; 4 G; 16 T; 0 other;

Query Match 2.7%; Score 26.8; DB 21; Length 30;  
 Best Local Similarity 93.3%; Pred. No. 1.3e+03; Mismatches 0; Indels 0; Gaps 0;  
 Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 289 caagatggtaaaactttaaaaacaaact 318  
 Db 30 CAGATGGACAAAGTTAAAACAAAGACT 1

RESULT 4  
 ID AAA93190 standard; DNA; 30 BP.  
 XX  
 AC AAA93190;  
 DT 11-JAN-2001 (first entry)

XX *Campylobacter jejuni* interrogation probe 11450.  
 KW *Campylobacter jejuni*; nucleic acid detection; genomic typing;  
 KW mutation detection; viral load determination; species identification;  
 KW forensic analysis; probe; ss.  
 OS *Campylobacter jejuni*.  
 XX  
 PN WO200049179-A1.

PD 24-AUG-2000.  
 XX  
 PF 18-FEB-2000; 2000WO-US04176.  
 XX  
 PR 18-FEB-1999; 99US-02252436.  
 PR 21-JUL-1999; 99US-0358972.  
 PR 27-SEP-1999; 99US-0406147.  
 XX  
 PA (PROM-) PROMEGA CORP.  
 XX  
 PT Shultz JW, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;  
 PI Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;  
 DR WPI; 2000-549282/50.

PT Detecting the presence of predetermined exogenous nucleic acid target  
 PT sequence useful for e.g. genotyping, comprises depolymerizing the 3'  
 PT end of an oligonucleotide probe hybridized to a nucleic acid target  
 PT sequence .  
 XX  
 PS Claim 47; Page 187; 230pp; English.

XX  
 CC The present sequence is an interrogation probe which was used to detect a  
 CC segment of the genome of *Campylobacter jejuni*. This was performed as part  
 CC of a method for determining the presence of a known exogenous nucleic  
 CC acid target sequence in a nucleic acid sample. The method comprises  
 CC adding a treated sample with a depolymerising enzyme which releases one  
 CC or more nucleotides from the 3'-end of a hybridised nucleic acid probe.  
 CC The method is used for assaying nucleic acids for a particular native or  
 CC mutant sequence, and for genomic typing. It is useful for detecting  
 CC mutations, translocations, and single nucleotide polymorphisms.  
 CC determination of viral load, species identification, detection of sample  
 CC contamination, and analysis of forensic samples. Compared with previous  
 CC methods of detecting nucleic acid hybrids, the new method has higher  
 CC sensitivity without the need for radiochemicals or electrophoresis. It is  
 CC quantitatively, highly reproducible and can be automated. The method can  
 CC reliably detect as few as 10 copies of a virus in a sample, and is  
 CC capable of providing multiple analyses in a single assay (multiplex  
 XX assay).  
 Sequence 30 BP; 16 A; 4 C; 5 G; 5 T; 0 other;

Query Match 2.7%; Score 26.8; DB 21; Length 30;  
 Best Local Similarity 93.3%; Pred. No. 1.3e+03; Mismatches 0; Indels 0; Gaps 0;  
 Matches 28; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 289 caagatggtaaaactttaaaaacaaact 318  
 Db 1 caagatggacaaactttaaaaacaaact 30

RESULT 5  
 ID AAX2169/c  
 XX  
 AC AAX2169;  
 XX  
 DT 18-JUN-1999 (first entry)

XX Synthetic plasmid synlux4 construction oligonucleotide R53.  
 KW DNA Plasmid; lux A; lux B; *Vibrio fisheri*; luciferase; promoter;  
 KW tnt9 kanamycin/neomycin phosphotransferase; DNA synthesis;  
 KW replication competent double-stranded polynucleotide; ss.  
 OS Synthetic.  
 XX  
 PN WO914318-A1.  
 XX  
 PD 25-MAR-1999.  
 XX  
 PR 16-SEP-1998; 98WO-US19312.  
 PR 16-SEP-1997; 97US-0059017.  
 XX  
 PA (TEXA ) UNIV TEXAS SYSTEM.  
 PI Evans GA;  
 XX  
 DR WPI; 1998-244029/20.

XX  
 PT Synthesis of replication competent double-stranded polynucleotides  
 XX  
 PS Example 4; Fig 5E; 135pp; English.  
 XX  
 CC AAX2021-212 represent oligonucleotide primers that were used to  
 CC construct a synthetic DNA plasmid sequence synlux4, to demonstrate the  
 CC method of the invention. Within the synlux4 sequence are included the  
 CC sequences of lux A, lux B, the A and B components of the *Vibrio fisheri*  
 CC luciferase sequence, positions of pUC19 including the origin of  
 CC replication and replication stability sequences, and the promoter and  
 CC coding sequence for tnt9 kanamycin/neomycin phosphotransferase. The  
 CC specification describes a method for the synthesis of replication



ID AAA97417  
 XX AAA97417 standard; DNA; 54 BP.  
 AC XX  
 XX AAA97417;  
 DT 29-JAN-2001 (first entry)  
 XX  
 DE Pea wild-type *pra2* gene light-repressible promoter oligonucleotide, wt3.  
 XX GTP-binding protein *pra2*; pea; light repressible promoter;  
 KW photoinhibitory; expression cassette; transgenic plant;  
 KW deterioration prevention; storage; ss.  
 XX OS  
 Pisum sativum.  
 XX PN WO20055313-A1.  
 XX PD 21-SEP-2000.  
 XX PP 03-MAR-2000; 2000WO-JP01269.  
 XX PR 12-MAR-1999; 99JP-0066551.  
 PA (SUNR ) SUNTORY LTD.  
 XX PI Guerry P, Trust T, Burg E, Lee L;  
 DR WPI; 2000-376214/32.  
 XX PS Disclosure; Page 7; 43PP; English.  
 XX CC The *flaA* gene encodes the major flagellin subunit of the *Campylobacter*  
 CC coli flagellar filament. Part of the *flaA* polypeptide may be fused with  
 CC the maltose binding protein of *Escherichia coli* to make a recombinant  
 CC protein. When this protein is introduced into a host an immunological  
 CC response is triggered. Therefore the recombinant protein may be used as  
 CC a vaccine to protect against *C. coli* intestinal colonisation and the  
 CC diarrhoea it causes. This vaccine system is useful as it can  
 CC prevent the development of Guillain-Barre syndrome (GBS) which is seen  
 CC with whole cell *Campylobacter* vaccines. The present sequence is the  
 CC *flaA* 11 PCR primer that was used to amplify part of the *flaA* gene.  
 XX SQ Sequence 27 BP; 12 A; 6 C; 3 G; 6 T; 0 other;  
 CS Query Match 2.1%; Score 21; DB 21; Length 27;  
 CC Best Local Similarity 100.0%; Pred. No. 3.1e+04;  
 CC Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 CC QY 1 attacacacaatgtgcagca 21  
 CC Db 7 attacacacaatgtgcagca 27  
 RESULT 10  
 ID AAA27149/C  
 XX ID AAA27149 standard; DNA; 33 BP.  
 AC XX  
 AC AAA27149;  
 XX DT 11-SEP-2000 (first entry)  
 XX DE *Campylobacter* coli *flaA* gene primer *flaA-2*.  
 XX KW Flagellin; *flaA*; diarrhoea; Guillain-Barre syndrome;  
 KW vaccine; GBS; PCR primer; ss.  
 OS XX  
 Campylobacter coli.  
 XX PN WO200027205-A1.  
 XX PD 18-MAY-2000.  
 XX PP 12-NOV-1999; 99WO-US27195.  
 XX PR 12-NOV-1999; 99US-0108114.  
 XX PA (USAT ) US SEC.  
 XX PI Guerry P, Trust T, Burg E, Lee L;  
 DE XX  
 DE *Campylobacter* coli *flaA* gene primer *flaA-11*.  
 XX KW Flagellin; *flaA*; diarrhoea; Guillain-Barre syndrome;  
 KW vaccine; GBS; PCR primer; ss.  
 XX DR WPI; 2000-376214/32.

PT Campylobacter FlaA protein and coding sequence, useful in reducing  
 PT Campylobacter intestinal colonization  
 XX disclosure; Page 7; 43pp; English.

CC The flaA gene encodes the major flagellin subunit of the Campylobacter  
 CC coil flagellar filament. Part of the FlaA polypeptide may be fused with  
 CC the maltose binding protein of *Escherichia coli* to make a recombinant  
 CC protein. When this protein is introduced into a host an immunological  
 CC response is triggered. Therefore the recombinant protein may be used as  
 CC a vaccine to protect against *C. coli* intestinal colonisation and the  
 CC diarrhoea it causes. This vaccine system is useful as it can  
 prevent the development of Guillain-Barre syndrome (GBS) which is seen  
 CC with whole cell Campylobacter vaccines. The present sequence is the  
 CC flaA-2 PCR primer that was used to amplify part of the flaA gene.  
 XX Sequence 33 BP; 10 A; 8 C; 1 G; 14 T; 0 other;

Query Match 2.1%; Score 21; DB 21; Length 33;  
 Best Local Similarity 100.0%; Pred. No. 3.3e+04; Indels 0; Gaps 0;  
 Matches 21; Conservative 0; Mismatches 0; CC

OY 979 gttaaaaatgtatgtttagat 999  
 |||||||  
 33 GTTAAATGTATGTTAGAT 13

Db

RESULT 11  
 ID AAT60508/C  
 XX AAT60508 standard; DNA; 24 BP.  
 AC  
 XX AAT60508;  
 DT 10-JUN-1997 (first entry)  
 XX DE Primer CFO4R.4.  
 XX KW PCR; polymerase chain reaction; amplify; infection; forensic science;  
 KW infectious pathogen; genetic disorder; genetic variance; primer; ss;  
 OS Synthetic.  
 XX PN US5612473-A.  
 XX PR 18-MAR-1997.  
 XX PF 16-JAN-1996; 96US-0587209.  
 XX PR 16-JAN-1996; 96US-0587209.  
 XX PA (GULL-) GULL LAB.  
 XX PT Coombs J, Glass MJ, Malmstrom SL, Wu L;  
 XX DR WPI; 1997-192163/17.

XX Processing samples for amplification of nucleic acid target  
 PT sequences - using extraction buffer containing at least one  
 PT detergent and a salt composition of greater than 1 molar  
 PT concentration  
 PS Example 3; Column 17; 21pp; English.

CC AAT60503-T60514 represent amplification primers for DNA sequences  
 CC present in a sample processed by the method of the invention. The  
 CC processing method of the invention comprises obtaining a sample of  
 CC material potentially containing the target nucleic acid sequences, and  
 CC mixing the sample with an external buffer solution. The buffer solution  
 CC comprises two detergents, and at least one salt composition present in a  
 CC greater than 1 M concentration. The mixture then centrifuged to obtain  
 CC a supernatant portion, which is then heated before being reconstituted  
 CC to precipitate the proteins, and obtaining a second supernatant portion,  
 CC

Query Match 2.1%; Score 20.8; DB 19; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3.4e+04; Indels 0; Gaps 0;  
 Matches 22; Conservative 0; Mismatches 2; CC

CC Sequence 24 BP; 6 A; 5 C; 5 G; 8 T; 0 other;  
 SQ

RESULT 12  
 ID AAV31446/C  
 XX AAV31446 standard; DNA; 24 BP.  
 AC  
 XX AAV31446;  
 DT 11-AUG-1998 (first entry)  
 XX DE Campylobacter nucleic acid sequence amplifying primer CFO4R.  
 XX KW Salmonella; microorganism; detection; multiple analyte; PCR primer;  
 KW Yersinia; *Escherichia coli*; Campylobacter; ss.  
 XX OS Synthetic.  
 OS Campylobacter sp.  
 XX PN US5756701-A.  
 XX PD 26-MAY-1998.  
 XX PR 06-AUG-1996; 96US-0592725.  
 XX PR 16-JAN-1996; 96US-0587209.  
 XX PR 06-AUG-1996; 96US-0592725.  
 XX PA (GULL-) GULL LAB INC.  
 XX PI Coombs J, Glass MJ, Malmstrom SL, Wu L;  
 XX DR WPI; 1998-321634/28.

XX PT Nucleic acid probes and primers - for detecting *Salmonella*, *Yersinia*  
 PT or *E. coli*  
 XX Claim 5; Column 17; 21pp; English.  
 XX  
 CC This primer is used for the PCR amplification of Campylobacter nucleic  
 CC acid sequences. The invention provides nucleic acid probes and primers  
 CC for detecting *Salmonella*, *Yersinia* or *E. coli*. It provides methods and  
 CC apparatus for detecting and discriminating multiple analytes within a  
 CC test sample. The methods are simple, user-friendly, cost effective and  
 CC fast. The methods and the probes and primer sequences are used for  
 CC detecting the corresponding microorganisms in clinical samples.  
 XX Sequence 24 BP; 6 A; 5 C; 5 G; 8 T; 0 other;

Query Match 2.1%; Score 20.8; DB 19; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3.4e+04; Indels 0; Gaps 0;  
 Matches 22; Conservative 0; Mismatches 2; CC

CC from which nucleic acids are precipitated. The isolated nucleic acids  
 CC are then dissolved. The method provides a rapid means of preparing a  
 CC sample for amplification so that multiple analytes can be detected and  
 CC differentiated within a relatively short time period (typically less  
 CC than 5 hours with the novel pre-processing step taking less than 5  
 minutes). Typical applications of nucleic acid amplification include  
 CC detection of infections in patients, foodstuffs and for  
 CC diagnostic/forensic or quality control purposes, to discriminate between  
 CC multiple potential infectious pathogens, to diagnose genetic disorders or  
 CC to identify genetic variances.  
 XX Sequence 24 BP; 6 A; 5 C; 5 G; 8 T; 0 other;

Query Match 2.1%; Score 20.8; DB 18; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3.4e+04; Indels 0; Gaps 0;  
 Matches 22; Conservative 0; Mismatches 2; CC

OY 91 ggcttttggatcaactccgcaca 114  
 |||||||  
 Db 24 GGCTTGGATCAACTCAGCAGCA 1

QY 91 ggctttagaaatcaactccggcaga 114  
 Db 24 GGCTTAACTAACAGCAGCA 1  
 Db 24 GGCTTAACTAACAGCAGCA 1

RESULT 13  
 AAV2594 2/C  
 ID AAV2594 standard; DNA: 24 BP.  
 XX  
 AAV2594:2;  
 XX  
 DT 15-JUL-1998 (first entry)  
 XX  
 DE Oligonucleotide PCR primer CFO4R gene.  
 XX  
 KW Sequence-specific probe; enterohaemorrhagic; Escherichia coli;  
 KW salmonella; Campylobacter; Shigella; Yersinia; beta-globin;  
 KW gastroenteritis; PCR primer; ss.  
 OS Synthetic.  
 OS Campylobacter sp.  
 XX  
 PN US575344-A.  
 XX  
 PD 19-MAY-1998.  
 XX  
 PF 07-AUG-1996; 96US-0689235.  
 XX  
 PR 16-JAN-1996; 96US-0587209.  
 PR 07-AUG-1996; 96US-0689235.  
 XX  
 PA (GULL-) GULL LAB INC.  
 XX  
 PT Coombs J, Glass MJ, Malmstrom SL, Wu L;  
 XR DR WPI; 1998-311393/27.

XX  
 PT Distinguishing between similar nucleic acid samples - using  
 PT sequence-specific probes e.g. between enterohaemorrhagic and normal  
 F\* Escherichia coli

XX  
 PS Example 3; Column 17; 21pp; English.

The present sequence represents a PCR primer used in an example of the present invention. The present invention describes a method for detecting mismatches between first and second nucleic acid sequences having at least one base difference. The method comprises: (a) obtaining at least one labelled probe consisting of an oligonucleotide sequence spanning the location of at least one base difference between the first and second sequences, where the oligonucleotide sequence contains at least one neutral base molecule in a position other than the position of the base difference(s) but is otherwise exactly complementary to the first sequence, so that the probe hybridises more weakly with the second sequence than with the first sequence; (b) mixing the probe(s) with the first and second sequences under hybridisation conditions; (c) dissociating any probe/second sequence hybrids; and (d) detecting probe/first sequence hybrids. The method can be used to distinguish between similar DNA/RNA sequences in a sample, especially to distinguish stool samples from people suffering from gastroenteritis, caused specifically by enterohaemorrhagic E. coli. The use of the method shortens the time between sample preparation to obtaining results, than has been possible with previous similar procedures.

XX  
 SQ Sequence 24 BP; 6 A; 5 C; 5 G; 8 T; 0 other;

Query Match 2.1%; Score 20.8; DB 19; Length 24;  
 Best Local Similarity 91.7%; Pred No. 3.4e+04; Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 Matches 22; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 91 ggctttagaaatcaactccggcaga 114  
 Db 24 GGCTTAACTAACAGCAGCA 1  
 XX  
 RESULT 15  
 AAF1671L  
 ID AAF1671 standard; DNA: 44 BP.  
 XX  
 AAF1671;  
 AC  
 XX  
 DT 09-MAR-2001 (first entry)

DE dgMMP-specific aptamer clone #19.

Polymeric biomolecules: aptamer: ss

Synthetic

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{PKAE-} PKAELUX INC.

Kwagh J, Macklin JJ, Mitsis RG, Ulmer KM;

WPI; 2001-016410/02. -

### Sequencing a polymer

polysaccharide or polypeptide, comprises separating a terminal monomer from the polymeric biomolecule and identifying the separated terminal

卷之三

Claim 3; Fig 13; 123pp; English.

The present invention relates to polymeric biocatalysts, the method

monomer from the polymeric biomol-

sequencing or structurally charac-

method is also useful for develop-

**Sequence** 44 BP; 9 A; 6 C; 19 G; 1

Very Match 2.18; Score 70.0%  
at Local Similarity

tches 28; Conservative 0; M

747 tgctatcaatgggtttagtaag

4 tgacaccactgggtgggtatgggtagg

ch completed: April 17, 2002, 02:17:16

time: 3512 sec

sequencing a polymeric biomolecule, such as a polynucleotide, polysaccharide or polypeptide, comprises separating a terminal monomer from the polymeric biomolecule and identifying the separated terminal monomer using an aptamer -

Claim 37; Fig 15; 123pp; English.

The present invention relates to a new method for sequencing a polymeric biomolecule. The method involves separating a terminal monomer from the polymeric biomolecule and identifying the separated terminal monomer using an aptamer. The method is useful for sequencing or structurally characterizing a polymeric biomolecule such as a polynucleotide, a polysaccharide or a polypeptide. The method is also useful for developing aptamers.

Sequence 44 BP; 9 A; 6 C; 19 G; 10 T; 0 other;

Query Match 2.1%; Score 20.8; DB 22; Length 44;

Query	Match	Score	Length	
Best	Local	Similarity	Score	Length
Y	tgctatcaatgggttataggtaaqtgttattcca	20.8	22	
4	tgcacccatgggttgttatggtaggttgttggaaata	3.9e+04	44	
Matches	28; Conservative	0; Mismatches	12; Indels	0; Gaps

SEARCH CENTERED. 2004-12-3003 03.12.16

Job time: 3512 sec





MEDIUM TYPE: Diskette, 3.50 inch,  
 MEDIUM TYPE: 1.44 Mb storage  
 COMPUTER: IBM compatible  
 OPERATING SYSTEM: MS-DOS  
 SOFTWARE: WordPerfect 6.0a for WINDOWS  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/587,209  
 FILING DATE: 16-JAN-1996  
 CLASSIFICATION: 435  
 INFORMATION FOR SEQ ID NO: 6:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 24 base Pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA  
 US-08-587-209-6

Query Match 2.1%; Score 20.8; DB 1; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3e+03; 0; Mismatches  
 Matches 22; Conservative 0; Indels 0; Gaps 0;  
 QY 91 ggcttttagaaatcaactccgcaga 114  
 ||||||| ||||| ||||| |||||  
 Db 24 GGCTTAGATAACTCGCAGCA 1

RESULT 6  
 Sequence 6, Application US/08689236  
 Patent No. 5736995  
 GENERAL INFORMATION:  
 APPLICANT: Wu, Linxian  
 APPLICANT: Coombs, Jana  
 APPLICANT: Malmstrom, Sharon L.  
 APPLICANT: Glass, Michael J.  
 APPLICANT: Coombs, Jana  
 APPLICANT: Malmstrom, Sharon L.  
 APPLICANT: Glass, Michael J.  
 TITLE OF INVENTION: Methods and Apparatus for Preparing, Amplifying, and Discriminating Multiple Analytes  
 NUMBER OF SEQUENCES: 30  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: David O. Seeley, Esq.  
 ADDRESSEE: Workman, Nydegger & Seeley  
 STREET: 1000 Eagle Gate Tower  
 STREET: 60 East South Temple  
 CITY: Salt Lake City  
 STATE: Utah USA  
 COUNTRY: USA  
 ZIP: 84111

COMPUTER READABLE FORM:  
 MEDIUM TYPE: Diskette, 3.50 inch,  
 MEDIUM TYPE: 1.44 Mb storage  
 COMPUTER: IBM compatible  
 OPERATING SYSTEM: MS-DOS  
 SOFTWARE: WordPerfect 6.0a for WINDOWS  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/689,235  
 FILING DATE: 16-JAN-1996  
 CLASSIFICATION: 435  
 INFORMATION FOR SEQ ID NO: 6:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 24 base Pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA  
 US-08-689-235-6

RESULT 7  
 Sequence 6, Application US/08689235  
 Patent No. 575601  
 GENERAL INFORMATION:  
 APPLICANT: Wu, Linxian  
 APPLICANT: Coombs, Jana  
 APPLICANT: Malmstrom, Sharon L.  
 APPLICANT: Glass, Michael J.  
 APPLICANT: Coombs, Jana  
 APPLICANT: Malmstrom, Sharon L.  
 APPLICANT: Glass, Michael J.  
 TITLE OF INVENTION: Methods and Apparatus for Preparing, Amplifying, and Discriminating Multiple Analytes  
 NUMBER OF SEQUENCES: 30  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: David O. Seeley, Esq.  
 ADDRESSEE: Workman, Nydegger & Seeley  
 STREET: 1000 Eagle Gate Tower  
 STREET: 60 East South Temple  
 CITY: Salt Lake City  
 STATE: Utah USA  
 COUNTRY: USA  
 ZIP: 84111

COMPUTER READABLE FORM:  
 MEDIUM TYPE: Diskette, 3.50 inch,  
 MEDIUM TYPE: 1.44 Mb storage  
 COMPUTER: IBM compatible  
 OPERATING SYSTEM: MS-DOS  
 SOFTWARE: WordPerfect 6.0a for WINDOWS  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/689,235  
 FILING DATE: 16-JAN-1996  
 CLASSIFICATION: 435  
 INFORMATION FOR SEQ ID NO: 6:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 24 base Pairs  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA  
 US-08-689-235-6

RESULT 8  
 Sequence 6, Application US/08692725  
 Patent No. 575601  
 GENERAL INFORMATION:  
 APPLICANT: Wu, Linxian  
 APPLICANT: Coombs, Jana  
 APPLICANT: Malmstrom, Sharon L.  
 APPLICANT: Glass, Michael J.  
 APPLICANT: Glass, Michael J.  
 TITLE OF INVENTION: Methods and Apparatus for Preparing, Amplifying, and Discriminating Multiple Analytes  
 NUMBER OF SEQUENCES: 30  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: David O. Seeley, Esq.  
 ADDRESSEE: Workman, Nydegger & Seeley  
 STREET: 1000 Eagle Gate Tower  
 STREET: 60 East South Temple  
 CITY: Salt Lake City

Query Match 2.1%; Score 20.8; DB 1; Length 24;  
 Best Local Similarity 91.7%; Pred. No. 3e+03; 0; Mismatches  
 Matches 22; Conservative 0; Indels 0; Gaps 0;

RESULT 9  
 Sequence 6, Application US/08692725  
 Patent No. 575601  
 GENERAL INFORMATION:  
 APPLICANT: Wu, Linxian  
 APPLICANT: Coombs, Jana  
 APPLICANT: Malmstrom, Sharon L.  
 APPLICANT: Glass, Michael J.  
 APPLICANT: Glass, Michael J.  
 TITLE OF INVENTION: Methods and Apparatus for Preparing, Amplifying, and Discriminating Multiple Analytes  
 NUMBER OF SEQUENCES: 30  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: David O. Seeley, Esq.  
 ADDRESSEE: Workman, Nydegger & Seeley  
 STREET: 1000 Eagle Gate Tower  
 STREET: 60 East South Temple  
 CITY: Salt Lake City



TITLE OF INVENTION: DIFFERENTIATION-SUPPRESSIVE POLYPEPTIDE  
FILE REFERENCE: KP-8576  
CURRENT APPLICATION NUMBER: US/09/214,278  
CURRENT FILING DATE: 1999-01-26  
NUMBER OF SEQ ID NOS: 32  
SOFTWARE: Patentin ver. 2.1

SEQ ID NO: 30  
LENGTH: 58  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Description of Artificial Sequence: synthetic DNA

US-09-214-278-30

RESULT 12  
US 08-886-967-3/C  
Sequence 3, Application US/08886967  
; Patent No. 606893  
GENERAL INFORMATION:  
APPLICANT: ASTOLFI, SPARTACO  
APPLICANT: DE LIMA, BEATRIZ D.  
APPLICANT: THIEMANN, JOSEF E.  
APPLICANT: TUNES DE SOUSA, HELOISA R.  
APPLICANT: VILELA, LUCIANO  
TITLE OF INVENTION: VECTOR FOR EXPRESSION OF HETEROLOGOUS PROTEIN AND METHODS FOR EXTRACTING RECOMBINANT PROTEIN AND FOR PURIFYING ISOLATED RECOMBINANT INSULIN

NUMBER OF SEQUENCES: 7  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: FROMMER LAWRENCE & HAUG LLP  
STREET: 745 FIFTH AVENUE  
CITY: NEW YORK  
STATE: NEW YORK  
COUNTRY: USA

ZIP: 10151  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/306,949  
CLASSIFICATION:  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 08/886,967  
ATTORNEY/AGENT INFORMATION:  
NAME: HAUG, EDGAR H.  
REGISTRATION NUMBER: 29,309  
REFERENCE/DOCKET NUMBER: 540519-2003  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 212-588-0800  
TELEFAX: 212-588-0500

INFORMATION FOR SEQ ID NO: 3:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 52 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)

US-09-306-949-3

RESULT 14  
US-08-874-825-118/C  
Sequence 118, Application US/08874825  
; Patent No. 6057101  
GENERAL INFORMATION:  
APPLICANT: Rothberg, Jonathan  
; APPLICANT: Nandabalan, Krishnan

Query Match 2.1%; Score 20.6; DB 3; Length 52;  
Best Local Similarity 67.4%; Pred. No. 4.6e+03;  
US-08-886-967-3

Query Match 2.1%; Score 20.6; DB 4; Length 58;  
Best Local Similarity 67.4%; Pred. No. 4.3e+03;  
US-09-306-949-3/C

Qy 400 ttaatgttgttttaccaatcaagaatccaaatcggtcaa 442  
; Sequence 3, Application US/09306949  
; Patent No. 62811329  
; GENERAL INFORMATION:  
; APPLICANT: ASTOLFI, SPARTACO  
; APPLICANT: DE LIMA, BEATRIZ D.  
; APPLICANT: THIEMANN, JOSEF E.  
; APPLICANT: TUNES DE SOUSA, HELOISA R.  
; TITLE OF INVENTION: VECTOR FOR EXPRESSION OF HETEROLOGOUS PROTEIN AND METHODS FOR EXTRACTING RECOMBINANT PROTEIN AND FOR PURIFYING ISOLATED RECOMBINANT INSULIN  
; NUMBER OF SEQUENCES: 7  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: FROMMER LAWRENCE & HAUG LLP  
; STREET: 745 FIFTH AVENUE  
; CITY: NEW YORK  
; STATE: NEW YORK  
; COUNTRY: USA  
; ZIP: 10151  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patentin Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/306,949  
; CLASSIFICATION:  
; PRIORITY APPLICATION DATA:  
; APPLICATION NUMBER: US 08/886,967  
; ATTORNEY/AGENT INFORMATION:  
; NAME: HAUG, EDGAR H.  
; REGISTRATION NUMBER: 29,309  
; REFERENCE/DOCKET NUMBER: 540519-2003  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 212-588-0800  
; TELEFAX: 212-588-0500  
; INFORMATION FOR SEQ ID NO: 3:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 52 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)

Qy 400 ttaatgttgttttaccaatcaagaatccaaatcggtcaa 442  
; Sequence 118, Application US/08874825  
; Patent No. 6057101  
; GENERAL INFORMATION:  
; APPLICANT: Rothberg, Jonathan  
; APPLICANT: Nandabalan, Krishnan

APPLICANT: Yang, Meijia  
 APPLICANT: Knight, James  
 APPLICANT: Kalbfleisch, Theodore  
 TITLE OF INVENTION: IDENTIFICATION AND COMPARISON OF PROTEIN-PROTEIN INTERACTIONS THAT OCCUR IN POPULATIONS  
 TITLE OF INVENTION: PROTEIN-PROTEIN INTERACTIONS THAT OCCUR IN POPULATIONS  
 NUMBER OF SEQUENCES: 122  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Penile & Edmonds  
 STREET: 1155 Avenue of the Americas  
 CITY: New York  
 STATE: NY  
 COUNTRY: USA  
 ZIP: 10036/2711  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Diskette  
 COMPUTER: IBM Compatible  
 OPERATING SYSTEM: DOS  
 SOFTWARE: FastSEQ Version 2.0  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/874,825  
 FILING DATE: 13-JUN-1997  
 CLASSIFICATION: 435  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: 08/653,824  
 FILING DATE: 14-JUN-1996  
 ATTORNEY/AGENT INFORMATION:  
 NAME: Mistock, S. Leslie  
 REFERENCE/DOCKET NUMBER: 7934-045  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: 212-790-9090  
 TELEFAX: 212-899-8864  
 TELEX: 66141 PENNIE  
 INFORMATION FOR SEQ ID NO: 118:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 39 base pairs  
 SEQUENCE:  
 TYPE: nucleic acid  
 STRANDEDNESS: single  
 TOPOLOGY: linear  
 MOLECULE TYPE: DNA  
 FEATURE:  
 NAME/KEY: misc-feature  
 LOCATION: 1..50  
 OTHER INFORMATION: /product= "OLIGOMER FOR  
 OTHER INFORMATION: CONSTRUCTION OF SYNTHETIC LD78 GENE"  
 US-08-450-905B-7

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Query Match 2.0%; Score 20; DB 3; Length 39;  
 Best Local Similarity 82.1%; Pred. No. 6e+03;  
 Matches 23; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

Qy 768 aggtaaatgttggatattcagatgtat 795  
 Db 39 AGGTCAGGTGCTTCAGATGTCAT 12

RESULT 15  
 US 08-450-905B-7/c  
 Sequence 7, Application US/08450905B  
 Patent No. 585601  
 GENERAL INFORMATION:  
 APPLICANT:  
 TITLE OF INVENTION: Stem Cell Inhibiting Proteins  
 NUMBER OF SEQUENCES: 178  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: HALE and DORR  
 STREET: 60 State Street  
 CITY: Boston  
 STATE: MA  
 ZIP: 02109  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 SOFTWARE: PatentIn Release #1.0, Version #1.25  
 CURRENT APPLICATION DATA:

Query Match 2.0%; Score 20; DB 2; Length 50;  
 Best Local Similarity 72.2%; Pred. No. 6.6e+03;  
 Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;  
 Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 368 caataccaccttcattttatggcaacaactttaa 403  
 Db 36 CAAATTCCACAAATTCTATTCGCTGACTTGTAA 1

Search completed: April 17, 2002, 02:18:56  
 Job time: 3297 sec

~ wed apr 17 07:36:45 2002

us-09-439-311-1.rni

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GenCore version 4.5  
copyright (C) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 17, 2002, 01:16:14 ; Search time 1464.96 Seconds  
(without alignments)  
7327.868 Million cell updates/sec

Title:	US-09-439-311-1
Perfect score:	999
Sequence:	1attaatacataatgttgcagc.....ttaaaatgtatggtagagat 999
scoring table:	IDENTITY_NUC Gapop 10.0 , Gapext 1.0
Searched:	11151937 seqs, 5372889281 residues
Total number of hits satisfying chosen parameters:	111874
Minimum DB seq length:	0
Maximum DB seq length:	60
Post-processing:	Minimum Match 0%
	Maximum Match 100%
Database :	Listing first 45 summaries
1:	EST:*
2:	em_estfun:*
3:	em_estin:*
4:	em_eston:*
5:	em_lespl:*
6:	em_esiba:*
7:	em_estro:*
8:	em_lescov:*
9:	em_htc:*
10:	gb_est1:*
11:	gb_est2:*
12:	gb_htc:*
13:	gb_gss:*
14:	em_gss_fun:*
15:	em_gss_hum:*
16:	em_gss_inv:*
17:	em_gss_pn:*
18:	em_gss_pro:*
19:	em_gss_rod:*
20:	em_gss_vrt:*
21:	em_gss_other:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

### SUMMARIES

Result No.	Score	Query Length	DB ID	Description
C 1	23.6	2.4	50	AU104183
C 2	21.5	2.2	60	AU104183
C 3	21.0	2.1	52	AU104183
C 4	20.8	2.1	59	AU104183
C 5	20.8	2.1	60	AU104183
C 6	20.6	2.1	58	AU104183
C 7	20.4	2.0	51	AU104183
C 8	20.4	2.0	55	AU104183
C 9	20.2	2.0	50	AU104183
C 10	20.2	2.0	54	AU104183
C 11	20.2	2.0	58	AU104183
C 12	20.2	2.0	59	AU104183

### ALIGNMENTS

RESULT	REFERENCE	AUTHORS	DEFINITION	ACCESSION	VERSION	KEYWORDS	SOURCE	ORGANISM
1	AU101183/c		AU104183 Sugano Homo sapiens mRNA sequence.	KAT06093,				
				AU104183	1	EST:13553704		
				AU104183		EST.		
						human.		
						Homo sapiens		
						Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
						(bases 1 to 50)		
						Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hatakeyama,T., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubou,K., Sugaya,A. and Sugano,S.		
						Fine structural analysis of transcription start sites of human mRNAs using full-length enriched and 5'-end enriched cDNA libraries		
						Unpublished (2001)		
						Contact: Yutaka Suzuki		
						Department of Virology		
						Institute of Medical Science, University of Tokyo		
						4-6-1, Shirokanedai, Minato-ku, Tokyo 108-8639, Japan		
						Email: yusukieims.u-tokyo.ac.jp		
						Suzuki,Y., Ioshimoto-Nakagawa,K., Maruyama,K., Sugaya,A. and Sugano,S.		
						Construction and characterization of a full length-enriched and a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).		
						FEATURES source		
						Location/Qualifiers		
						1. -50		
						/organism="Homo sapiens"		
						/db_xref="taxon:9606"		
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						ORIGIN	4 g	22 t





FEATURES		Location/Qualifiers
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1. .58		musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource ( <a href="http://www.Jax.org/resources/documents/dnarecs/">http://www.Jax.org/resources/documents/dnarecs/</a> ). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptored DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD2 (9147321149b1AF29072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptored mouse DNA was annealed to adaptored vector DNA, and transformed into chemically-competent E. coli XL1-Gold (Stratagene) cells and selected for ampicillin resistance.
Query Match	2.1%	Score 20.6; DB 10; Length 58;
Best Local Similarity	62.7%	Pred. No. 7.7e+05;
Matches	32;	Mismatches 19; Indels 0; Gaps 0;
OY	208	atcttgcactcagaataaggcatgtatggcaacttaaacttagat 258
ACCESSION	AZ812517	
VERSION	AZ812517.1	GI:12981841
KEYWORDS		
SOURCE		
ORGANISM		
RESULT	7	
LOCUS	AZ812517	51 bp DNA
DEFINITION	2M0079C15F	Mouse 10kb plasmid UGCGCM library Mus musculus genomic
ACCESSION		
VERSION		
KEYWORDS		
SOURCE		
ORGANISM		
REFERENCE		
AUTHORS	Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A., and Weiss, D., Weiss, R.	
TITLE		Mouse whole genome scaffolding with paired end reads from 10kb plasmid insert
JOURNAL		Unpublished (2000)
COMMENT		Contact: Robert B. Weiss University of Utah Genome Center Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA Tel: 801 505 5606 Fax: 801 505 7177 Email: <a href="mailto:ddunn@genetics.utah.edu">ddunn@genetics.utah.edu</a>
FEATURES		
source		
1. .51		High quality sequence stop: 51.
FEATURES		
source		Location/Qualifiers
1. .51		/organism="Mus musculus" /strain="C57BL/6J" /clone="UUGCGCM0079C15F" /lab_xref="taxon:10090" /sex="Female" /sex="Male" /lab_host="E. coli strain XL10-gold, T1-resistant, F-" /note="vector: pMD2uv; Purified genomic DNA from M."
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ORIGIN		
BASE COUNT	9 a	18 c 11 g 13 t
ORIGIN		
RESULT	8	
LOCUS	AU014315	55 bp mRNA
DEFINITION	AU014315	Schizosaccharomyces pombe late log phase cDNA
ACCESSION	AU014315	Schizosaccharomyces pombe cDNA clone spcc09537, mRNA sequence.
VERSION	AU014315.1	GI:3369106
KEYWORDS		
SOURCE		
ORGANISM		
REFERENCE		
AUTHORS	Morimoto, M. and Mita, K.	
TITLE		Identification of expressed sequence tags of Schizosaccharomyces pombe
JOURNAL		Unpublished (1998)
COMMENT		Contact: Mitsuaki Morimoto Genome Research Group National Institute of Radiological Sciences 9-1, Anagawa-4-chome, Inage-ku, Chiba, Chiba 263-8555, Japan Email: <a href="mailto:morimotol@rs.nirs.go.jp">morimotol@rs.nirs.go.jp</a>
FEATURES		
source		
1. .55		Location/Qualifiers
FEATURES		
source		
1. .51		/organism="Schizosaccharomyces pombe" /strain="972" /clone="spcc09537" /clone_1b="Schizosaccharomyces pombe late log phase cDNA" /sex="h minus" /note="vector: M13mp19; the cDNA library of Schizosaccharomyces pombe was prepared by cloning cDNA into the small site of M13mp19 DNA and the direction of DNA sequences was not always from 5' to 3'. The cDNA data of Schizosaccharomyces pombe are available for searching on the World Wide Web. (URL, <a href="http://www.nirs.go.jp">http://www.nirs.go.jp</a> )"
BASE COUNT	24 a	2 c 9 g 20 t
ORIGIN		
Query Match	2.0%	Score 20.4; DB 10; Length 55;

Qy	172	ggtcaggatatactaaaggatgtatgtttggatcttgcaa	217	Best Local Similarity 65.2%; Pred NO. 8.6e+05; Matches 30; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
Db	2	GGTTATGTTAATAGTATAGTAAACGATTATGTATGTGCTAA	47	Accession AU103731.1 GI:13553252
COMMENT				RESULT 9 AU103731/C AU103731 50 bp mRNA EST 05-APR-2001 LOCUS AU103731 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone DEFINITION COL04386, mRNA sequence. ACCESSION AU103731 VERSION AU103731.1 KEYWORDS EST. TITLE Homo sapiens ORGANISM Eukaryota; Metazoa; Chordata; Craniata; vertebrata; Euteleostomi; Mammalia; Butheria; Primates; Catarrhini; Hominidae; Homo. REFERENCE 1 (bases 1 to 50) AUTHORS Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata H., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo K., Sugaya,A. and Sugano,S. TITLE Fine structural analysis of transcription start sites of human mRNAs using full-length enriched and 5'-end enriched cDNA libraries JOURNAL Unpublished (2001) COMMENT Contact: Yutaka Suzuki Department of Virology Institute of Medical Science, University of Tokyo 4-6-1, Shirokanedai, Minato-ku, Tokyo 108-8639, Japan Email: yuzuki@ims.u-tokyo.ac.jp Suzuki,Y., Yoshihimo-Nakabu,K., Maruyama,K., Sugaya,A. and Sugano S., Construction and characterization of a full length-enriched and a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).
FEATURES	Source			High quality sequence stop: 54. Location/Qualifiers 1..54 /organism="Mus musculus" /strain="C57BL/6J" /db_xref="taxon:10900" /clone="UGCC2M1220G15" /sex="Female" /lab_host=E. coli strain XL10-Gold, T1-resistant, F-" /note="vector: PWD4Inv; purified genomic DNA from M. musculus C57BL/6J (female) was obtained from the Jackson Laboratory Mouse DNA Resource ( <a href="http://wwwjax.org/resources/documents/dnars/">http://wwwjax.org/resources/documents/dnars/</a> ). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (gi 473214 gb AF129072.1), a copy-number inducible derivative of pBluescript. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptors vector DNA, and transformed into chemically competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."
BASE COUNT	6 a 12 c 13 g 19 t	ORIGIN	18 a 9 c 12 g 15 t	RESULT 10 Query Match 2.0%; Score 20.2; DB 10; Length 50; Best Local Similarity 63.3%; Pred. No. 9.6e+05; Matches 31; Conservative 0; Mismatches 18; Indels 0; Gaps 0; Qy 636 ttgcgttggatcacacagaatgcgatataacacaaatcgatcgca 684 Db 50 TTGGCGCAGGAAGCGACCAACATGGACAGAACACAATCCACGCA 2
COMMENT				RESULT 10 Query Match 2.0%; Score 20.2; DB 10; Length 50; Best Local Similarity 63.3%; Pred. No. 9.6e+05; Matches 31; Conservative 0; Mismatches 18; Indels 0; Gaps 0; Qy 636 ttgcgttggatcacacagaatgcgatataacacaaatcgatcgca 684 Db 50 TTGGCGCAGGAAGCGACCAACATGGACAGAACACAATCCACGCA 2
BASE COUNT	18 a 9 c 12 g 15 t	ORIGIN	18 a 9 c 12 g 15 t	RESULT 11 Query Match 2.0%; Score 20.2; DB 13; Length 54; Best Local Similarity 63.3%; Pred. No. 9.6e+05; Matches 31; Conservative 0; Mismatches 18; Indels 0; Gaps 0; Qy 846 tgttcaggccatctaaggatggaaatgtaaactgttcttacttcgccc 894 Db 6 TGTCCTAAGTGTAAGATGAGAATAGGAACCTGCCTCTCTTACACC 54
FEATURES	Source			RESULT 11 BF131272/C LOCUS BF131272 58 bp mRNA EST 24-OCT-2000 DEFINITION 6010195281 NIH_MGC_58 Homo sapiens cDNA clone IMAGE:4051302 5', mRNA sequence. ACCESSION AU295454 VERSION AU295454.1 GI:13825771 KEYWORDS GSS. TITLE house mouse. SOURCE house mouse. ORGANISM Mus musculus COMMENT Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Butheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. Dunn,D., Aoyagi,A., Barber,M., Beacons,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenah,E., Pedersen,T., Reilly M., Rose,M., Rose,R., Stokes,R., Tingay,A., von Niederhausern,A. and Wright,D., Weiss,R. TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
REFERENCE	1 (bases 1 to 54)	REFERENCE	1 (bases 1 to 58)	COMMENT Unpublished (2000) Contract: Robert B. Weiss University of Utah Genome Center University of Utah, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT Rm. 308, Biomedica
AUTHORS	Dunn,D., Aoyagi,A., Barber,M., Beacons,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenah,E., Pedersen,T., Reilly M., Rose,M., Rose,R., Stokes,R., Tingay,A., von Niederhausern,A. and Wright,D., Weiss,R.	AUTHORS NIH-MGC <a href="http://mgc.nci.nih.gov/">http://mgc.nci.nih.gov/</a> . TITLE National Institutes of Health, Molecular Gene Collection (MGC). JOURNAL Unpublished (1999) COMMENT Contact: Robert Strausberg, Ph.D. Email: <a href="mailto:cabpsbr@mail.nih.gov">cabpsbr@mail.nih.gov</a>		

FEATURES	source
source	<p>Tissue Procurement: ATCC      cDNA library Preparation: CLONETECH Laboratories, Inc.      CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)      DNA Sequencing by: Incyte Genomics, Inc.      Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:  <a href="http://image.llnl.gov">http://image.llnl.gov</a>      Plate: LICH886 row: ] column: 07.</p>
Location/Qualifiers	<p>Location/Qualifiers</p>
1..58	<p>1..58</p>
/organism="Homo sapiens"	<p>/organism="Homo sapiens"</p>
/lab_xref="txxon_9606"	<p>/lab_xref="txxon_9606"</p>
/clone="IMAGE:4051302"	<p>/clone="IMAGE:4051302"</p>
/clone_1ib="NIR_MGC_58"	<p>/clone_1ib="NIR_MGC_58"</p>
/tissue_type="hypernephroma"	<p>/tissue_type="hypernephroma"</p>
/lab_host="DHL10B (T1 phage-resistant)"	<p>/lab_host="DHL10B (T1 phage-resistant)"</p>
/note="Organ: kidney; Vector: PDRN-LIB (clonitech); Site_1: SII (ggccgcgtcgcc); Site_2: SII (ggcccttatgcgc); Double-stranded cDNA was prepared from cell line RNA. 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CACGCCATATGCC-3', and 3' adaptor sequence: 5'-ATTTAGGGCCAGGGGCCGAT-3')BN-3', (where B = A, C, or G and N = A, C, G, or T) Average insert size 1.35 kb (range 0.9-4.0 kb). 15/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech laboratories (Palo Alto, CA)."	
BASE COUNT	<p>22 a 9 c 0 g 26 t 1 others</p>
ORIGIN	<p>ORIGIN</p>
Query Match	<p>Query Match</p>
Best Local Similarity 63.3%	<p>Score 20.2; DB 11; Length 58;</p>
'Matches 31; Conservative 0; Mismatches 18; Indels 0; Gaps 0;	
Qy	<p>543 tgtaggacttacataaaacctacaacgttatcgatggattttaaattt 501</p>
Db	<p>49 TGATAATAGATAATTAATTAAATGGATGTAAGGATGAAAT 1</p>
RESULT 12	<p>RESULT 12</p>
W38842/c	<p>W38842</p>
LOCUS	<p>W38842</p>
DEFINITION	<p>59 bp mRNA</p>
IMAGE:3049105, similar to gb:M90516	<p>EST</p>
VERSION	<p>15-MAY-1996</p>
KEYWORDS	<p>Homo sapiens</p>
SOURCE	<p>IMAGE:3049105</p>
ORGANISM	<p>Human.</p>
ACCESSION	<p>W38842</p>
VERSION	<p>W38842.1 GI:1320547</p>
REFERENCE	<p>Mammalia; Butheria; Chordata; Craniata; Vertebrata; Euteleostomi; VERBENACEAE</p>
AUTHORS	<p>Miller,L., Clark,N., Dubugue,T., Elliston,K., Hawkins,M., Holman,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J., Rikfin,L., Rohlfing,T., Soares,M., Tan,F., Treviskis,E., Waterston,R., Williamson,A., Wohldmann,P. and Wilson,R.</p>
TITLE	<p>The WashU-Merck EST Project</p>
JOURNAL	<p>Unpublished (1995)</p>
COMMENT	<p>Contact: Wilson RR      Washington University School of Medicine      4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108      Tel: 314 286 1800      Fax: 314 286 1810      Email: estwatson.wustl.edu      IMAGE Consortium (<a href="http://image.llnl.gov">http://image.llnl.gov</a>) for further information.      Trace considered overall poor quality      Seq primer: mob REGA-ET      High quality sequence stop: 1.</p>
FEATURES	<p>FEATURES</p>
source	<p>source</p>
/organism="Mus musculus"	<p>/organism="Mus musculus"</p>
1..59	<p>1..59</p>
/organism="Homo sapiens"	<p>/organism="Homo sapiens"</p>
/ab_xref="GDB:1248320"	<p>/ab_xref="GDB:1248320"</p>
/db_xref="taxon:9606"	<p>/db_xref="taxon:9606"</p>
/clone="IMAGE:304910"	<p>/clone="IMAGE:304910"</p>
/clone_1ib="Soares-parathyroid_tumor_NbHPA"	<p>/clone_1ib="Soares-parathyroid_tumor_NbHPA"</p>
/tissue_type="parathyroid tumor"	<p>/tissue_type="parathyroid tumor"</p>
/dev_stage="adult"	<p>/dev_stage="adult"</p>
/lab_host="DHL10B (ampicillin resistant)"	<p>/lab_host="DHL10B (ampicillin resistant)"</p>
/note="Organ: parathyroid gland; Vector: PR773D (Pharmacia ) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a NOT I - Oligo(dT) primer	<p>/note="Organ: parathyroid gland; Vector: PR773D (Pharmacia ) with a modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA was primed with a NOT I - Oligo(dT) primer</p>
/trypT_3]	<p>/trypT_3]</p>
Qy	<p>550 cttaacttataaaactacacgttatcgaaatggatgtg 598</p>
Db	<p>51 CTGTCCTCTAAATCACAATCACAGACCTGCTATTAAATCCATATG 3</p>
RESULT 13	<p>RESULT 13</p>
AZ434413	<p>AZ434413</p>
LOCUS	<p>AZ434413</p>
DEFINITION	<p>54 bp DNA</p>
VERSION	<p>IM0220118R Mouse 10kb plasmid UGGCIM library Mus musculus genomic clone UGGCIM0220118 R, DNA sequence.</p>
ACCESSION	<p>AZ434413</p>
KEYWORDS	<p>house mouse</p>
SOURCE	<p>GSS</p>
ORGANISM	<p>Organism</p>
REFERENCE	<p>Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.</p>
AUTHORS	<p>Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duvall,B., Hamil,C., Islam,H., Longcore,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A., and Wright,D. Weiss,R.</p>
TITLE	<p>Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts</p>
JOURNAL	<p>Unpublished (2000)</p>
COMMENT	<p>Contact: Robert B. Weiss      University of Utah Genome Center      University of Utah      Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA      Tel: 801 585 5606      Fax: 801 585 7177      Email: ddunn@genetics.utah.edu      Insert Length: 10000 Std Error: 0.00</p>
Plate: 0220 row: I column: 18	<p>Plate: 0220 row: I column: 18</p>
Seq primer: CACCAAGGAAACAGCTATGACC	<p>Seq primer: CACCAAGGAAACAGCTATGACC</p>
Class: plasmid ends	<p>Class: plasmid ends</p>
High Quality sequence stop: 54	<p>High Quality sequence stop: 54</p>
1..54	<p>1..54</p>
Location/Qualifiers	<p>Location/Qualifiers</p>
FEATURES	<p>FEATURES</p>
source	<p>source</p>

FEATURES		vector to vector length is 57.
source		Location/Qualifiers
/clone_libr="Mouse 10kb plasmid UGCGCIM0220118"		1. .56
/clone_libr="Mouse 10kb plasmid UGCGCIM0220118"		/organism="Glycine max"
/sex="Male"		/db_xref="taxon:3847"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"		/clone_libr="Gm-c1037"
/note="Vector: PWD2nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource		/tissue, type="fully expanded leaves of greenhouse grown plants", dev_stage="2 week old"
(http://www.Jax.org/resources/documents/dnases/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptored DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD2 (g11473214gb1_AF129072.1), a copy-number inducible derivative of plasmid RL. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptored mouse DNA was annealed to adaptored vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."		/lab_host="DH10B"
ORIGIN		/note="Vector: psprt1; site_1: NotI; site_2: SalI; This cDNA library was constructed from mRNA isolated from fully expanded leaves of greenhouse grown plants that were 2 weeks old. The library was prepared using the Life Technologies Psuperscript cDNA library construction kit. Complementary DNA was synthesized from mRNA using a poly(dT) sequence with a NotI restriction site. SalI linkers adaptors were ligated to the blunt-ended cDNA fragments followed by NotI digestion. The cDNA fragments were directionally cloned into the NotI-SalI restriction site of the pREPORT vector. The ligated cDNA fragments were transformed into E. coli Electro- Max DH10B host cells. This library was constructed in the laboratory of Dr. Lila Vodkin at the University of Illinois at Urbana-Champaign. email: l-vodkin@uic.edu"
BASE COUNT		18 t
19 a		11 c
6 g		18 t
14		18 t
RESULT		Query Match 2.0%; Score 20; DB 13; Length 54;
LOCUS		Best Local Similarity 61.5%; Pred. No. 1.1e+06;
DEFINITION		Matches 32; Conservative 0; Mismatches 0; Gaps 0;
Gm-c1037-803 5', mRNA sequence.		Indels 0; Version AWT80772
ACCESSION		EST: AWT80772.1 GI:7705447
VERSION		
KEYWORDS		
SOURCE		
ORGANISM		
Glycine max		Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophytina		Rosidae; eurosids I; Fabales; Fabaceae; Papilionoideae; Phaseoleae;
Glycine		Glycine; Glycine
1 (bases 1 to 56)		(bases 1 to 56)
REFERENCE		
AUTHORS		Shoemaker,R., Keim,P., Vodkin,L., Erpelding,J., Coryell,V., Khanna,A., Boila,B., Maria,M., Hillier,L., Kucaba,T., Martin,T., Beck,C., Wylie,T., Underwood,K., Stepien,M., Theising,B., Allen,M., Bowers,Y., Person,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,P., Waterston,R., Wilson,R.
TITLE		Public Soybean EST Project
JOURNAL		Unpublished (1999)
COMMENT		Contact: Shoemaker R/Public Soybean EST Project
COMMENT		Washington University School of Medicine
COMMENT		4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
COMMENT		Tel: 314 286 1800
COMMENT		Fax: 314 286 1810
COMMENT		Email: est@watson.wustl.edu
COMMENT		This clone is available through: Genome Systems, Inc. 4633 World Parkway Circle St. Louis, Missouri 63134 For further information call: (800) 430-0030 or (314) 247-3322 FAX: (888) 919-3324 or (314) 427-3324 or contact: clones@genomesystems.com or info@genomesystems.com web site: www.genomesystems.com Putative full length read
BASE COUNT		18 t
15 a		2 c
11 g		28 t
BASE COUNT		28 t
ORIGIN		
RESULT		Query Match 2.0%; Score 20; DB 10; Length 56;
LOCUS		Best Local Similarity 65.9%; Pred. No. 1.1e+06;
DEFINITION		Matches 29; Conservative 0; Mismatches 15; Indels 0; Gaps 0;
GSS		Version AFI49647 Human chromosome 18q21 from exon-trapping Homo sapiens
ACCESSION		AF149647
VERSION		AF149647.1 GI:8485985
KEYWORDS		GSS,
SOURCE		human.
ORGANISM		Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
1 (bases 1 to 56)		
REFERENCE		
AUTHORS		Chen,H., Huo,Y., Patel,S., Zhu,X., Swift-Scanlan,T., Reeves,R.H., DePaulo,R.Jr., Ross,C.A. and McInnis,M.G.
TITLE		Gene identification using exon amplification on human chromosome 18q21: implications for bipolar disorder
JOURNAL		Mol. Psychiatry 5 (5), 502-509 (2000)
MEDLINE		2005132
COMMENT		Contact: Chen H
COMMENT		Psychiatry and Behavioral Sciences
COMMENT		Johns Hopkins University School of Medicine
COMMENT		600 N. Wolfe Street, Baltimore, MD 21287, USA
COMMENT		Email: hc@welchlink.welch.jhu.edu
FEATURES		Location/Qualifiers
source		1. .56
/organism="Homo sapiens"		/db_xref="taxon:9606"
/map="18q21"		/clone_libr="Human chromosome 18q21 from exon-trapping"
BASE COUNT		13 t
24 a		12 c
7 g		13 t

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Query Match          2.0%; Score 20; DB 13; length 56;
Best Local Similarity 61.5%; Pred NO. 1.1e-06;
Matches 32; Conservative 0; Mismatches 20; Indels 0; Gaps 0;
Y 335 tcacccgtttatggaaactgtataatcccaaattaccacttcatttaa 386
  ||||| | | | | | | | | | | | | | | | | | | | | | | | | | | |
b 4 TCGAGCAGCCAAAGAAAAGATTTGTCATACCAACAGATGGCATCATTTA 55

```

Search completed: April 17, 2002, 02:14:13  
Job time: 3479 sec

OM of: US-09-439-311-2 to: N\_Geneseq\_1101;\* out\_format : pfs  
 Date: Apr 17, 2002 3:13 AM  
 About: Results were produced by the GenCore software, version 4.5,  
 Copyright: (c) 1993-2000 Compugen Ltd.

## Command line parameters:

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-MODEL=frame+P2N.model -DBV=xlp
-O/cgn2-1/uspmo.spool/US9939311/runat_16042002_134011_11729/app_query.fasta_1.395
-DB=N_Geneseq_1101 -QFMT=fasta -SUFFIX=mg -GAPOP=12.000
-GAPLEN=4.000 -MINMATCH=100 -LOOPBL=0.000 -LOOPEXT=0.000
-OGAPOP=.500 -OGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500
-FGAPOP=6.000 -FGAPEXT=7.000 -RGAPOP=10.000 -YGAPEXT=0.500
-DELOM=6.000 -DELEXTR=7.000 -START=1 -MATRIX=blossom2
-TRANS=human40.gcd -LIST=45 -DCALIGN=200 THR_SCORE=pct
-THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=pfs
-NCPU=6 -ICPU=3 -LONGLOG -NO_XLPXY -WAIT -THREADS=1
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## Search information block:

Query: US-09-439-311-2

Query length: 333

Database: N\_Geneseq\_1101;\*

Database sequences: 930621

Database length: 42862619

Search time (sec): 170.970000

## score\_list:

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2000.DAT:AAAB6891	-		50.00	100.69	1.6e+03	30

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2000.DAT:AAAB6892	+		50.00	100.69	1.6e+03	30

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2000.DAT:AAAB3188	-		50.00	100.69	1.6e+03	30

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1998.DAT:AAAB93190	+		50.00	100.69	1.6e+03	30

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1998.DAT:AAV23196	+		88.52	7.4e-03	59	L

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1999.DAT:AAV4813	+		43.00	85.32	1.1e+04	51

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1998.DAT:AAV00234	+		41.00	82.48	1.6e-04	50

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2000.DAT:AAZ66924	+		40.00	79.47	2.4e+04	59

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1995.DAT:AAV00254	+		40.00	79.32	2.4e+04	60

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2001.DAT:AAV70806	+		40.00	79.32	2.4e+04	60

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2001.DAT:AAV30690	+		39.00	83.80	1.4e+04	31

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1998.DAT:AAV56429	-		39.00	80.03	2.2e+04	47

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1999.DAT:AAV21884	-		39.00	80.03	2.2e+04	47

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1999.DAT:AAV69197	-		39.00	80.03	2.2e+04	47

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2000.DAT:AAH79171	-		39.00	80.03	2.2e+04	47

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2000.DAT:AAH79171	-		39.00	80.03	2.2e+04	47

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA2000.DAT:AAH79171	-		39.00	80.03	2.2e+04	47

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1995.DAT:AAV5069	-		38.00	77.43	3.1e-04	53

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1995.DAT:AAV5069	-		38.00	77.43	3.1e-04	53

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1995.DAT:AAV5069	-		38.00	76.46	3.5e+04	59

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
/SIDS2/gcdata/geneseq/geneseq/NA1995.DAT:AAV5069	-		38.00	76.31	3.5e+04	60

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/SIDS2/gcdata/geneseq/geneseq/NA2001.DAT:AAH79776 - 36.00 74.76 4.3e-04 51
/SIDS2/gcdata/geneseq/geneseq/NA2001.DAT:AAU66898 - 36.00 74.76 4.3e-04 51
/SIDS2/gcdata/geneseq/geneseq/NA2001.DAT:AAH38820 - 36.00 74.76 4.3e-04 51
/SIDS2/gcdata/geneseq/geneseq/NA1995.DAT:AAQ7415 + 36.00 74.41 4.5e-04 53
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seq\_name: /SIDS2/gcdata/geneseq/geneseq/NA2000.DAT:AAAB6891

seq\_documentation-block:

ID	AAA86891 standard; DNA; 30 BP.
XX	
AC	AAA86891;

15-JAN-2001 (first entry)

Probe to Campylobacter jejuni.

Detection; nucleic acid hybrid; depolymerisation; analysis; SNP; single nucleotide polymorphism; identification; primer; target; mutation; genotyping; medical marker diagnostic; primer; target; mutation; genetic disease; ss.

Campylobacter jejuni.

W0200049180-11.

24-AUG-2000.

XX

18-FEB-2000; 2000WO-US04242.

XX

PR 21-JUL-1999; 99US-0358972.

PR 25-AUG-1999; 99US-0383316.

XX (PROM-) PROMEGA CORP.

Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;

PI Schultz JW, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;

PT Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;

XX WPI: 2000-565377/52.

Determining presence or absence of a predetermined endogenous nucleic acid sequence by using an enzyme that depolymerizes the 3' end of an oligonucleotide probe hybridized to a target sequence to release identifier nucleotides

Example; Page 321; 389pp; English.

The present invention describes a method (M1) for determining the presence or absence of a predetermined endogenous nucleic acid target sequence (ENR). The method comprises hybridising a probe having an identifier nucleotide (IN) with ENR which is treated with an enzyme that depolymerises the 3' end of hybridised ENR to release the INs.

M1 is used for determining the number of known sequence repeats present in a nucleic acid target sequence in a nucleic acid sample. The method is also useful for determining whether a nucleic acid target sequence in a sample is an allele from a homozygous or heterozygous locus. The method is also useful for detection of mutations, translocations and SNPs in nucleic acids (including those associated with genetic disease), determination of viral load, species identification, sample contamination, and analysis of forensic samples. AA86791 to AAA87079 and AAB12817 represent sequence which are used in the exemplification of the present invention.

There is a discrepancy between the SEQ ID NO: and sequences given in the examples, and the SEQ ID NO: and sequences given in the sequence listing from the present invention.

Sequence 30 BP; 5 A; 5 C; 4 G; 16 T; 0 other;

alignment\_scores: Quality: 50.00 Length: 10 Ratio: 5.00 Gaps: 0

Percent Similarity: 100.00 Percent Identity: 100.00



X Sequence 30 BP; 5 A; 5 C; 4 G; 16 T; 0 other;  
 SQ alignment\_scores:  
 Quality: 50.00 Length: 10  
 Ratio: 5.000 Gaps: 0  
 Percent Similarity: 100.000 Percent Identity: 100.000  
 alignment\_block:  
 US-09-439-311-2 x AAA93188/rev ..  
 Align seg 1/1 to reverse of: AAA93188 from: 1 to: 30  
 97 GlnaspGlyGlnSerLeuLysThrArgTr 106  
 ||||||| ||||| ||||| ||||| |||||  
 30 CAAGATGGACAAGTTAAACAGAACT 1  
 seq\_name: /SIDS2/seqdata/geneseq/geneseq/NA2000.DAT:AAA93190  
 seq\_documentation\_block:  
 ID AAA93190 standard; DNA; 30 BP.  
 XX AC AAA93190:  
 XX DT 11-JAN-2001 (first entry)  
 DE Campylobacter jejuni interrogation probe 11450.  
 XX KW Campylobacter jejuni; nucleic acid detection; genomic typing;  
 KW mutation detection; viral load determination; species identification;  
 forensic analysis; probe; ss.  
 OS Campylobacter jejuni.  
 XX PN WO200049179-A1.  
 XX PD 24-AUG-2000.  
 XX PF 18-FEB-2000; 2000WO-US04176.  
 XX PR 18-FEB-1999; 99US-0282436.  
 PR 21-JUL-1999; 99US-0358972.  
 PR 27-SEP-1999; 99US-0406147.  
 XX PA (PROM-) PROMEGA CORP.  
 XX PI Shultz JW, Lewis MK, Leippe D, Mandrekar M, Kephart D, Rhodes RB;  
 PI Andrews CA, Hartnett JR, Gu T, Olson RJ, Wood KV, Welch R;  
 XX DR WPI: 2000-549282/50.  
 XX PT Detecting the presence of predetermined exogenous nucleic acid target  
 sequence useful for e.g. genotyping, comprises depolymerizing the 3'  
 end of an oligonucleotide probe hybridized to a nucleic acid target  
 sequence -  
 XX PS Claim 47; Page 187; 230pp; English.  
 XX The present sequence is an interrogation probe which was used to detect a  
 segment of the genome of Campylobacter jejuni. This was performed as part  
 of a method for determining the presence of a known exogenous nucleic  
 acid target sequence in a nucleic acid sample. The method comprises  
 admiring a treated sample with a depolymerising enzyme which releases one  
 or more nucleotides from the 3' end of a hybridised nucleic acid probe.  
 The method is used for assaying nucleic acids  
 mutant sequence, and for genomic typing. It is useful for detecting  
 mutations, translocations, and single nucleotide polymorphisms,  
 determination of viral load, species identification, detection of sample  
 contamination, and analysis of forensic samples. Compared with previous  
 methods of detecting nucleic acid hybrids, the new method has higher  
 sensitivity without the need for radiochemicals or electrophoresis. It is  
 quantitative, highly reproducible and can be automated. The method can  
 reliably detect as few as 10 copies of a virus in a sample, and is  
 capable of providing multiple analyses in a single assay (multiplex  
 assay).  
 CC sequence 30 BP; 16 A; 4 C; 5 G; 5 T; 0 other;  
 CC alignment\_scores:  
 Quality: 50.00 Length: 10  
 Ratio: 5.000 Gaps: 0  
 Percent Similarity: 100.000 Percent Identity: 100.000  
 alignment\_block:  
 US-09-439-311-2 x AAA93190 ..  
 Align seg 1/1 to: AAA93190 from: 1 to: 30  
 97 GlnaspGlyGlnSerLeuLysThrArgTr 106  
 ||||||| ||||| ||||| ||||| |||||  
 1 CAAGATGGACAAGTTAAACAGAACT 30  
 seq\_name: /SIDS2/seqdata/geneseq/geneseq/NA1998.DAT:AAV23196  
 seq\_documentation\_block:  
 ID AAV23196 standard; DNA; 59 BP.  
 XX AC AAV23196:  
 XX DT 28-JUL-1998 (first entry)  
 DE Lactococcus lactis constitutional promoter Cp29.  
 XX KW Lactococcus lactis; constitutional promoter; optimise; spacer;  
 KW artificial promoter library; gene expression; ds.  
 XX OS Synthetic.  
 OS Lactococcus lactis.  
 XX FH Key Location/Qualifiers  
 FT Promoter 4..59  
 FT /tag= a  
 FT /standard\_name= "Constitutional promoter"  
 XX PN WO9807846-A1.  
 XX PD 26-FEB-1998.  
 XX PF 25-AUG-1997; 97WO-DK00342.  
 XX PR 23-AUG-1996; 96DK-0000886.  
 XX PA (JENS/) JENSEN P R.  
 XX PI Hammer K, Jensen PR;  
 XX DR WPI: 1998-179062/16.  
 XX PT New artificial promoter libraries - containing consensus promoter  
 PT sequences and variable spacers, used to generate promoters for  
 PT optimising expression of genes  
 XX PS Claim 28; Page 52; 89pp; English.  
 XX This is a Lactococcus lactis constitutional promoter sequence used in the  
 construction of an artificial promoter library of the invention. The  
 artificial promoter library for a selected organism or group of organisms  
 comprise a mixture of double-stranded DNA fragments, the sense strands of  
 which comprise at least half of two consensus sequences of efficient  
 promoters from the organism or group of organisms and surrounding or  
 intermediate nucleotide sequences (spacers) of variable length in which  
 at least 7 nucleotides are selected randomly, with the proviso that  
 previously known promoter sequences and promoter sequences isolated from  
 natural sources are not included. This promoter library can be used in a

method of optimising the expression of a gene in a microorganism. The method comprises selecting a set of promoters covering a range of promoter activities in relatively small steps of activity change from such an artificial promoter library and cloning the set of promoters into the organism placing in each clone the gene under the control of at least one promoter from the set and growing the selected clones and screening them to find the one showing optimised flux of product formation. Promoters covering wide ranges of activities, including very strong promoters can be generated which can be used for optimising expression of genes.

Sequence 59 BP; 15 A; 9 C; 16 G; 19 T; 0 other;  
**SQ**

alignment_scores:	Quality:	Length:	Gaps:	Percent Identity:
	46.00	13	0	76.923

**Percent Similarity:** 76.923  
**alignment\_block:**  
**US-09-439-311-2 x AAV23196 ..**

**Align seg 1/1 to : AAV23196 from: 1 to: 59**

291 GlyLysLeuValLeuThrSerAlaAspGlyArgGlyIle 303  
 ||||| ||||| ||||| |||||  
 5 GGTAAAGTTATCTTGACATCTCAGGGGGACGTGCTATA 43

**seq\_name:** /STD\$2/gcdata/geneseq/geneseq/NA1999.DAT:AA\$34813

**seq\_documentation\_block:**  
**ID AAX34813; AC AAX34813;**

**DT 06-JUL-1999 (first entry)**

**DE Human ZSIG-11 DNA specific primer ZC13735.**

**KW Secretory protein; ZSIG-11; ligand polypeptide; testis; endoprotease; prohormone convertase; fertility; therapeutic; human; PCR primer; ss.**

**XX OS Synthetic. OS Homo sapiens.**

**PN WO9916870-A1.**

**PD 08-APR-1999.**

**PR 29-SEP-1998; 98WO-US20449.**

**PR 19-MAY-1998; 98US-0085966.**

**PR 29-SEP-1997; 97US-000327.**

**PR 19-MAY-1998; 98US-0081310.**

**DR 1999-263692/22.**

**XX PT Polynucleotide encoding a human secretory protein, ZSIG-11.**

**PS Example 9; Page 108; 113pp; English.**

The invention relates to a human secretory protein, ZSIG-11. Host cells containing a vector comprising the ZSIG-11 nucleic acid are used for the recombinant expression of the protein. ZSIG-11 is a novel ligand polypeptide and specific antibodies can be used to detect its presence in a biological sample. Probes derived from ZSIG-11 nucleotide sequences can also be used in detection of ZSIG-11 RNA. ZSIG-11 is expressed at high levels in testis, and could be used to identify/study prohormone

CC convertses or endoproteinases that exhibit testis specificity.  
 CC Antagonists, including antibodies, are useful for inhibiting or  
 CC eliminating the function of ZSIG-11. It is possible that ZSIG-11 and  
 CC its antagonists will be useful as fertility inducing therapeutics.  
 CC Sequences AAX34800-21 represent PCR primers for amplifying the ZSIG-11  
 CC DNA.  
**SQ Sequence 51 BP; 17 A; 5 C; 19 G; 10 T; 0 other;**

**alignment\_scores:**  
**Quality:** 43.00      **Length:** 16      **Gaps:** 0      **Percent Identity:** 50.000  
**Percent Similarity:** 75.000  
**alignment\_block:**  
**US-09-439-311-2 x AAX34813 ..**

**Align seg 1/1 to : AAX34813 from: 1 to: 51**

253 GlyValValIleGlyLysValAspTyrSerAspGlyAspGluAsnGly 268  
 ||||| :::: ||| |||||:::||| |||:::|||  
 1 GGTTGTAAGCTTGGACAAGAGAGATTACAGGACCATGATGACAGGGT 48

**seq\_name:** /STD\$2/gcdata/geneseq/geneseq/NA1999.DAT:AAX19519

**seq\_documentation\_block:**  
**ID AAX19519 standard; DNA; 51 BP.**

**XX AC AAX19519; XX DT 07-JUN-1999 (first entry)**

**DE Human lipocalin homologue zlipol PCR primer ZC13\_735.**

**KW Human; lipocalin; testis; mammary gland; breast tumour; zlipol; KW breast cancer; emphysema; skin disease; reproduction; anti-inflammatory; KW antimicrobial; PCR primer; ss.**

**XX OS Synthetic. OS Homo sapiens.**

**PN WO9907740-A2.**

**PD 18-FEB-1999.**

**XX PR 06-AUG-1998; 98WO-US16425.**

**XX PR 06-AUG-1997; 97US-0054867.**

**XX PA (ZYMO ) ZYMOGENETICS INC.**

**XX PI Conklin DC;**

**XX DR WPI; 1999-167367/14.**

**XX PT New lipocalin homologue designated zlipol - whose expression is restricted to testis and mammary gland tissues, particularly breast tumour tissue, used to, e.g. predict tumour aggressiveness.**

**XX PS Example 5; Page 89; 94pp; English.**

The present sequence represents a PCR primer for lipocalin homologue, zlipol. The lipocalin homologue, zlipol, is specifically expressed in testis and mammary gland, particularly breast tumour tissue. Based on this tissue distribution, zlipol may be used as a diagnostic for breast carcinomas and as a tool for predicting tumour aggressiveness. Agonists can be used for transportation of small hydrophobic molecules either *in vivo* or *in vitro*, and so are useful in specifically promoting the growth and/or development of testis-specific cell lineages in culture. Zlipol can be used to identify inhibitors. Zlipol proteins can also be used to prepare antibodies (which can be linked to toxins), and can serve as immunogens. Zlipol proteins can be used as a delivery and encapsulation

CC system to transport and/or stabilise small lipophilic molecules, e.g. to  
 CC protect from gut pH and digestive enzymes. They can also be used to bind  
 CC small fatty acids in blood or tissues to modulate their biological  
 CC function, e.g. to transport retinoids or steroids to receptors, in  
 CC particular as therapy for breast cancer, emphysema and diseases of the  
 CC skin. They may also play an important role in reproduction. Other uses  
 CC include anti-inflammatory responses, and antimicrobial activities.  
 CC zlipol nucleic acid sequences may be used for gene therapy to increase  
 CC inhibit zlipol activity to derive probes and primers, to derive  
 CC antisense sequences, and to detect genetic abnormalities.

Sequence 51 BP; 17 A; 5 C; 19 G; 10 T; 0 other;

alignement\_scores:

Quality:	43.00	Length:	16
Ratio:	3.583	Gaps:	0
Percent Similarity:	75.000	Percent Identity:	50.000

alignement\_block:  
 US-09-439-311-2 x AAX19519 ..

Align seg 1/1 to: AAX19519 from: 1 to: 51

253 GlyyalvallealleglylylvalasptyrSerAsparglyAspGluangly 268  
 ||||| ::||| ||||||| ||||| ||||:|||  
 1 GGTGTAAGCTTGACAGAGATGATGAGACATGAGCAGAGGT 48

seq\_name: /SIDS2/egedata/geneseq/geneseq/NA1998.DAT:AAV00234

seq\_documentation\_block:  
 ID AAV00234 standard; DNA; 50 BP.

AC AAV00234;

XX DT 08-JUN-1998 (first entry)

DE tick vasoactive amine binding protein FS-HBPI reverse PCR primer.

XX Female-specific vasoactive amine binding protein 1; FS-HCPI;  
 KW histamine; serotonin; assay; antihistamine; anti-inflammatory;

KW insect bite; snake bite; scorpion bite; dermatitis; vaccine;  
 transgenic animal; tick; PCR; primer; ss.

XX OS Synthetic.

OS Rhipephalus appendiculatus.

XX PN W09744451-A2.

XX PD 27-NOV-1997.

XX PF 19-MAY-1997; 97WO-GB01372.

XX PR 18-APR-1997; 97GB-0007844.

XX PR 18-MAY-1996; 96GB-0010484.

XX PA (OXFO-) OXFORD VACS LTD.

XX PI Nuttall PA, Paesen GC;

XX DR WPI; 1998-010506/02.

PT New vasoactive amine binding proteins and related nucleic acid,

PT vectors - transformed cells and transgenic animals, used for  
 PT assaying or removing histamine and as antihistamine or  
 PT anti-inflammatory agents

XX Example 3; Page 20: 44pp; English.

CC This reverse primer was used with a forward primer (see AAV00233)  
 CC to amplify the coding region (see AAV00227) of Rhipephalus  
 CC appendiculatus female-specific histamine binding protein 1  
 CC (FS-HBPI) (see AAV37446), a novel vasoactive amine binding protein

CC (VABP). The primers were designed so that a SacI site was added  
 CC upstream of the start codon, while the stop codon was replaced by  
 CC a BamHI site, followed by 6 histidine codons and an SpeI site  
 CC comprising a TAG stop codon. The PCR product was ligated into  
 CC transfer vector pACC1291, generating plasmid pACC1291-FSI-HIS.  
 CC FS-HBPI was expressed as a histidine-tagged protein in *Saccharomyces*  
 CC *fruiperda* SF21 ovarian cells using a baculovirus expression system.  
 CC VBPs can be used to assay or remove histamine, as an antihistamine  
 CC or anti-inflammatory agent, and in vaccines.

Sequence 50 BP; 11 A; 8 C; 14 G; 17 T; 0 other;

alignement\_scores:

Quality:	41.00	Length:	10
Ratio:	4.100	Gaps:	0
Percent Similarity:	100.000	Percent Identity:	70.000

alignement\_block:  
 US-09-439-311-2 x AAV00234 ..

Align seg 1/1 to: AAV00234 from: 1 to: 50

262 SerAspArglyAspGluAsnGlySerileuile 271  
 |||||||:|||||:|||||:|||||:|||:  
 7 AGTGAGTGTGATGATGGATCCCTTCCTG 36

seq\_name: /SIDS2/egedata/geneseq/geneseq/NA2000.DAT:AAZ96924

seq\_documentation\_block:  
 ID AAZ96924 standard; DNA; 59 BP.

AC AAZ96924;

XX DT 14-APR-2000 (first entry)

DE S. cerevisiae gene deletion cassette constructing primer YMR290c-S1.  
 XX PR Antimycotic; mycosis; immunodepression; AIBS; diabetes; fungicide;

KW mycete; gene deletion; PCR primer; ss.

XX OS Saccharomyces cerevisiae.

XX PN WO9955907-A2.

XX PD 04-NOV-1999.

XX PR 22-APR-1999; 99WO-EP02722.

XX PR 24-APR-1998; 98EP-0401007.

XX PR 11-SEP-1998; 98EP-0402254.

XX PA (HMR1) HOECHST MARTON ROUSSEL.

XX PI Diu-Hercend A, Entian K, Koetter P;

XX DR WPI; 2000-105527/09.

PT Identifying antimycotic substances useful for drug preparation and  
 PT treatment of mycosis -

XX PS Examples; Page 71; 86pp; English.

CC The invention provides a method of screening for antimycotic substances  
 CC using essential genes from mycetes or a functionally similar mycete  
 CC gene or the corresponding encoded protein as target. The essential gene  
 CC useful for screening antimycotic substances is selected from the  
 CC following genes: YML14C, YLR186W, YLR15C, YLR22C, YLR243W, YLR272C,  
 CC YLR275W, YLR276C, YLR373W, YLR359W, YLR342W, YLR437C, YLR40C,  
 CC YML023C, YML049C, YML077W, YML093W, YML127W, YMR032W, YMR093C, YMR131C,  
 CC YMR185W, YMR212C, YMR213W, YMR218C, YMR288W, YMR290C, YMR211W,  
 CC YMR49C, YMR134W, YDR293W, YDR365C, YDR396W, YDR407C, YDR216W,  
 CC YDR449C, YDR472W, YDR499W, YDR141C, YDR324C, YDR325W, YDR398W, YDR246W,

CC YDR36C, YDR361C, YDR367W, YDR339C, YDR429C, YDR468C, YDR494W,  
 CC YDR57W, YDR201W, YDR434W, YDR181C, YDR31W, YPL023W, YPL035W,  
 CC YPL038W, YPL024W, YPL020C, YPL012W, YPL007C, YPL146C, YIL091C,  
 CC YIL083C, YIL019W, YIL109C, YIL104C, YFL024C, YFR033C, YFR027W, YFR024W,  
 CC YIR010W, YIR015W, YPR048W, YPR072W, YPR082C, YPR085C, YPR105C, YPR112C,  
 CC YPR137W, YPR133W, YPR144C and YPR169W. The method is useful for  
 CC identifying substances for the preparation of drugs for the treatment of  
 CC mycosis or prevention in immunodepression states. Drugs containing  
 CC antimycotic substances are useful for the treatment of mycotic  
 CC infections which occur during diseases like AIDS or diabetes. Substances  
 CC which may be used for the fabrication of fungicides, especially of  
 CC fungicides which are harmless for humans and animals, and antimycotic  
 CC substances which selectively inhibit the growth of specific mycete  
 CC species only, can also be identified by this method. Sequences  
 CC AZ296811-966990 represent PCR primers used in construction of S.  
 CC cerevisiae deletion cassettes.

XX sequence 59 BP; 8 A; 12 C; 15 G; 24 T; 0 other;

SQ alignment\_scores:  
 Quality: 40.00 Length: 15  
 Ratio: 3.077 Gaps: 0  
 Percent Similarity: 86.667 Percent Identity: 53.333

alignment\_block:  
 US-09-439-311-2 x AAZ96924 ..

Align seg 1/1 to: AAZ96924 from: 1 to: 59

163 SerLysIleGlyVaThrArgPheGluGlySerGlnSerphe 177  
 9 ACGTCTTGGTATTGGCGTTTCACAGGCCAGCTGAAGCTC 53

seq\_name: /SIBS2/ggldata/geneseq/geneseq/NA1995.DAT:AAT00254

seq\_documentation\_block:  
 ID AAT00254 standard; DNA; 60 BP.

AC AAT00254;

XX DT 14-AUG-1996 (first entry)

XX DE Thrombin 60N DNA ligand, clone #31.

XX KW Family 1; family 2; ligand; thrombin;  
 KW systematic evolution of ligands by exponential enrichment; SELEX;  
 KW heparin; selection; region of homology; inhibitor; ss.

XX OS Synthetic.

XX PN WO9521853-A1.

XX PD 17-AUG-1995.

XX PF 06-FEB-1995; 95W0-US01458.

XX PR 28-MAR-1994; 94US-021012.

PR 10-FEB-1994; 94US-019505.

PR 11-JUN-1990; 90US-0536428.

PR 10-JUN-1991; 91US-0714131.

PR 22-APR-1993; 93US-0061691.

XX (NESS-) NEXSTAR PHARM INC.

XX PI Gold L, Janjic N, Tasset D;

XX DR WPI; 1995-293073/38.

XX PT Identification of ligands to basic fibroblast growth factor and  
 PT thrombin - which can be modified for increased in vivo stability

XX PS Claim 39; Page 97; 236pp; English.

XX sequence 60 BP; 10 A; 11 C; 29 G; 10 T; 0 other;

SQ alignment\_scores:  
 Quality: 40.00 Length: 19  
 Ratio: 2.667 Gaps: 0  
 Percent Similarity: 78.947 Percent Identity: 42.105

alignment\_block:  
 US-09-439-311-2 x AAT00254 ..

Align seg 1/1 to: AAT00254 from: 1 to: 60

204 ThrSerValGlyThrGlyLeuGlyLalaLeuAlaGluGluIleasnArgas 220  
 4 ACCGGGGAGGGCGTAGGGTTGGAGCGTGCCGATGTGGTAGGCACGGA 53  
 220 nalaAsp 222  
 ::::|||  
 54 CTCGGAT 60

seq\_name: /SIBS2/ggldata/geneseq/geneseq/NA2001.DAT:AAF70806

seq\_documentation\_block:  
 ID AAF70806 standard; DNA; 60 BP.

XX AC AAF70806;

XX DT 20-APR-2001 (first entry)

XX DE Thrombin high affinity ligand #53.

XX KW Ligand; basic fibroblast growth factor; bFGF; gene therapy; vascular;  
 KW atherosclerosis; angioplasty; stability; ss.

XX OS Unidentified.

XX PN US6177557-B1.

XX PD 23-JAN-2001.

XX PF 05-AUG-1996; 96US-0687421.

XX PR 11-JUN-1990; 90US-0536428.

PR 10-JUN-1991; 91US-0714131.

PR 06-NOV-1992; 92US-097333.

PR 10-FEB-1994; 94US-019505.

PR 28-MAR-1994; 94US-0219012.

XX (NESS-) NEXSTAR PHARM INC.

XX PI Janjic N, Gold L, Tasset D;

XX DR WPI; 2001-158583/16.

XX PT Novel nucleic acid ligands to basic fibroblast growth factor that are  
 PT useful as inhibitors of basic fibroblast growth factors and 2-amino  
 modified RNA ligands, exhibit increased in vivo stability



PT adhesion molecule R - with ligands HS-beta and tubulin using  
 PT two-hybrid assay, useful for treating inflammation, T cell  
 activation etc.

XX Example 13; Column 135-136; 108pp; English.

CC AAV56429-V56334 are primers used in the isolation of a novel human  
 CC intercellular adhesion molecule, ICAM-R. This sequence is used in a  
 CC method which investigates modulators of the interaction between ICAM-R  
 CC and the 14.3.3 family member HS-beta and tubulin. An anti-ICAM-R  
 CC antibody, can block, inhibit or stimulate ligand/receptor interactions  
 CC involving ICAM-R, particularly its effector functions involved in  
 CC (non)specific immune responses. ICAM-R related agents may be used to  
 CC treat or monitor inflammation, disorders involving T cell activation or  
 CC macrophages, e.g. adult respiratory distress syndrome, stroke, Crohn's  
 CC disease, multiple sclerosis, rheumatoid arthritis, asthma, tumour  
 CC growth, human immune deficiency virus infection, diabetes, graft vs. host  
 CC disease and many others. Antibodies may also be used for passive  
 CC immunisation, for purifying, detecting or quantifying ICAM-R and for  
 CC identifying ICAM-R expressing cells.

XX Sequence 47 BP; 9 A; 21 C; 7 G; 10 T; 0 other;

alignment\_scores:  
 Quality: 39.00 Length: 12  
 Ratio: 3.900 Gaps: 0  
 Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:  
 US-09-439-311-2 x AAV56429/rev ..

Align seg 1/1 to reverse of: AAV56429 from: 1 to: 47

seq\_name: /SIDS2/gcdata/geneseq/geneseqn/NA1999.DAT:AAV21884  
 seq\_documentation\_block:  
 ID AAV21884 standard; DNA; 47 BP.  
 XX  
 AC AAX21884;  
 XX  
 DT 14-MAY-1999 (first entry)

DE Primer for antibody against ICAM-R.

XX  
 KW ICAM: immunoglobulin-like loop; intercellular adhesion molecule receptor;  
 KW alpha d/CD18; antibody; immunisation; inflammatory response; asthma;  
 KW tumour growth; viral infection; therapy; primer; ss.

XX Synthetic.

OS Mus sp.

XX USS880268-A.

XX 09-MAR-1999.

PP 07-JUN-1995; 95US-0483932.

XX 05-AUG-1994; 94US-0286754.

PR 27-JAN-1992; 92US-0827689.

PR 26-MAY-1992; 92US-0889724.

PR 05-JUN-1992; 92US-0884061.

PR 22-JAN-1993; 93US-0009266.

PR 26-JAN-1993; 93W0-US00787.

PR 05-AUG-1993; 93US-0102852.

PR 07-JUN-1995; 95US-0483932.

PA (ICOS-) ICOS CORP.

XX Gallatin WM, Vazeux R;

XX WPT; 1989-204041/17.

DR

XX Example 13; Column 41; 108pp; English.

CC This sequence is a primer for DNA encoding an antibody specific for  
 CC ICAM-R. The invention relates to antibodies (Ab) which bind specifically  
 CC to the intercellular adhesion molecule receptor (ICAM-R), inhibiting the  
 CC interaction between ICAM-R and alpha d/CD18. Abs with specific ICAM-R  
 CC binding are useful in compositions for immunisation, and for purifying  
 CC ICAM-R polypeptides and identifying cells expressing ICAM-R on their cell  
 CC surface, modulating ligand/receptor binding and biological activities  
 CC involving ICAM-R, especially inflammatory responses of the specific  
 CC immune system, the non specific immune system, monitoring and treating  
 CC asthma, tumour growth, and/or metastasis, and viral infection (e.g. HIV  
 CC infection). In particular diseases involving an essential T cell  
 CC activation (e.g. asthma, psoriasis, diabetes, graft vs. host disease,  
 CC tissue transplant rejection, and multiple sclerosis) may be treated with  
 CC anti-ICAM-R antibodies. The Abs specifically bind to and identify ICAM-R  
 CC and disrupt ICAM-R to cell adhesion molecule, especially alpha d/CD18

XX SQ Sequence 47 BP; 9 A; 21 C; 7 G; 10 T; 0 other;

alignment\_scores:  
 Quality: 39.00 Length: 12  
 Ratio: 3.900 Gaps: 0  
 Percent Similarity: 83.333 Percent Identity: 66.667

alignment\_block:  
 US-09-439-311-2 x AAV21884/rev ..

Align seg 1/1 to reverse of: AAV21884 from: 1 to: 47

seq\_name: /SIDS2/gcdata/geneseq/geneseqn/NA1999.DAT:AAV69197  
 seq\_documentation\_block:  
 ID AAV69197 standard; DNA; 47 BP.  
 XX  
 AC AAV69197;  
 XX  
 DT 17-FEB-1999 (first entry)

DE Humanised ICR-1.1 antibody V<sub>k</sub> region DNA mutating oligo 110.

XX Intercellular adhesion molecule polypeptide; ICAM-R, humanised; ICR 1.1;

KW ICR 8.1; monoclonal antibody; therapeutic; inflammatory; asthma; tumour;  
 KW graft-versus-host disease; viral infection; toxin; radionuclide;  
 KW neovascularisation site; mutagenic; PCR primer; ss.

XX Synthetic.

OS Mus sp.

PN US5837822-A.

PP 17-NOV-1998.

XX 07-JUN-1995; 95US-0487113.

PR 07-JUN-1995; 95US-0487113.

PR 27-JAN-1992; 92US-0827689.

PR 26-MAY-1992; 92US-088972A.  
 PR 03-JUN-1992; 92US-0194061.  
 PR 22-JAN-1993; 93US-0009265.  
 PR 26-JAN-1993; 93WO-US00781.  
 PR 05-AUG-1993; 93US-0102852.  
 XX  
 PA (ICOS-) ICOS CORP.  
 XX  
 PT Gallatin WM, Vazeux R;  
 XX  
 WPI; 1999-023535/02.

Humanised antibodies specific for intercellular adhesion molecule polypeptide - useful for therapeutic or diagnostic purposes

Example 13; Column 42; 116pp; English.

XX  
 CC The invention relates to humanised ICR 1.1 and ICR 8.1 antibodies targeted to the human intercellular adhesion molecule polypeptide (ICAM-R) polypeptide. Antibodies specific for ICAM's are potentially useful as therapeutic compounds for treating e.g. immune-mediated inflammatory conditions (e.g. graft-versus-host disease), asthma, tumours or viral infections. Monoclonal antibodies specific for ICAM-R, or their conjugates formed with e.g. toxins or radionuclides are useful for therapeutically targeting or detecting neovascularisation sites. PCR mutagenic oligos AAV69197 and AAV69198 are used in the construction of the VK region of the humanised antibody ICR-1.1.  
 XX  
 SQ Sequence 47 BP; 9 A; 21 C; 7 G; 10 T; 0 other;

alignment\_scores:  
 Quality: 39.00 Length: 12  
 Ratio: 3.900 Gaps: 0  
 Percent Similarity: 83.333 Percent Identity: 66.667  
 alignment\_block:  
 05-09-439-311-2 x AAV69197/rev

Align seg 1/1 to reverse of: AAV69197 from: 1 to: 47

169 Arg Phe Glu Thr Gly Ser Gln Ser Phe Ser Gly 180

|||:||||||| |||:||||||| 44 AGGAATGGAGACTGGTCAGCACCGATTGGGAGTGA 9





J., DeFord,J., McFarland,J., Burzinski,K., Khan,M., Kupfer,K. and Garner,H.R.  
 Genomic Sequence Sampled Map of Chromosome 11  
 Unpublished (1996)  
 Contact: Evans GA, Shane Probst  
 McDermott Center for Human Growth and Development  
 University of Texas Southwestern Medical Center At Dallas  
 5323 Harry Hines Blvd, Dallas TX 75235-8591  
 Tel: 214-688-1600  
 Fax: 214-688-1666  
 Email: gevans@utsouthwestern.edu, shane@mcdermott.swmed.edu  
 Seq primer: T7  
 Class: cosmid ends  
 High quality sequence stop: 60.  
 Location/Qualifiers  
 FEATURES source  
 1. .60  
 /organism="Homo sapiens"  
 /clone="CSR1-2591"  
 /obj\_xref="taxon:606"  
 /clone.lib="CSRL flow sorted Chromosome 11 specific  
 cosmid"  
 /sex="female"  
 /cell\_type="chimeric hamster somatic cell hybrid"  
 /note="vector: scos-1; Human Chromosome 11 specific cosmid  
 library prepared from flow sorted human Chromosome 11  
 derived from Chinese Hamster Ovary (CHO) monochromosomal  
 somatic cell hybrid, J1."  
 BASE COUNT ORIGIN  
 16 a 16 c 5 g 22 t 1 others  
 Align seg 1/1 to: B04096 from: 1 to: 60  
 seq.name: gb\_gss:AZ469793  
 seq\_documentation\_block:  
 LOCUS AZ469793 44 bp DNA  
 DEFINITION 1M0203F04R Mouse 10kb plasmid UGGC1M library Mus musculus genomic  
 ACCESSION AZ469793  
 VERSION AZ469793.1 GI:10627918  
 KEYWORDS GSS.  
 SOURCE house mouse.  
 ORGANISM Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Butheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 REFERENCE 1 (bases 1 to 44)  
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Dongare,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A., and Wright,D., Weiss,R.  
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb  
 Plasmid inserts  
 JOURNAL Unpublished (2000)  
 COMMENT Contact: Robert B. Weiss  
 University of Utah Genome Center  
 ← University of Utah

---

Rn. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT  
 84112, USA  
 Tel: 801 585 5606  
 Fax: 801 585 7177  
 Email: ddunn@genetics.utah.edu  
 Insert Length: 10000 Std Error: 0.00  
 Plate: 0283 row: F column: 04  
 Seq primer: CACAGGAAACGCTAIGACC  
 Class: plasmid ends  
 High quality sequence stop: 44.  
 Location/Qualifiers  
 FEATURES source  
 1. .44  
 /organism="Mus musculus"  
 /strain="C57BL/6J"  
 /obj\_xref="taxon:6090"  
 /clone="UUGC1M0283F04"  
 /clone.lib="Mouse 10kb plasmid UGGC1M library"  
 /sex="Male"  
 /lab\_host="E. Coli strain XL10-Gold, T1-resistant, F-"  
 /note="Vector: PMD4 zny; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.Jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptored DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD2 (gll1472114.gbl|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptored mouse DNA was annealed to chemically competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance." (Stratagene) cells  
 BASE COUNT ORIGIN  
 9 a 12 c 12 g 11 t  
 Align seg 1/1 to: AZ469793 from: 1 to: 44  
 seq.name: gb\_est1:AU107968  
 seq\_documentation\_block:  
 LOCUS AU107968 50 bp mRNA  
 DEFINITION AU107968 Sugano Homo sapiens cDNA library Homo sapiens EST  
 ACCESSION AU107968  
 VERSION AU107968.1 GI:13557490  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo Sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Butheria; Primates; Catarhini; Hominidae; Homo;  
 KAT11118, mRNA sequence.  
 REFERENCE 1 (bases 1 to 50)  
 AUTHORS Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata,H., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo,K., Suyama,A. and Sugano,S.



```

/cultivar="mixed background W23/A188/B73"
/db_xref="taxon:4577"
/clone_1ib="006 - Rescuemu Grid G"
/tissue_type="leaf"
/dev_stage="adult"
/lab_host="DH10B"
/note="Organ: leaf; Vector: Rescuemu (engineered from pBluescript backbone); Site,1: BamHI; Site,2: BglII;
Rescuemu is a 4.9 kb, modified maize Mu transposon designed to allow plasmid rescue from total genomic DNA. Mu elements insert preferentially into transcription units. For more information on Rescuemu, go to the web site, www.zmud.lastate.edu' and follow the links for Rescuemu. Grid G was grown at Stanford in 2000. DNA was extracted from leaf punches, double digested using BamHI and BglII, and ligated to form circular plasmids. DH10B cells were transformed and then screened on LB plates with ampicillin."
BASE COUNT          13 a   18 c   7 g   19 t
ORIGIN

alignment_scores:
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    Ratio: 4.333        Gaps: 0
Percent Similarity: 81.818      Percent Identity: 54.545
alignment_block:
Align seg 1/1 to reverse of: AZ21603  from: 1 to: 57
US-09-439-311-2 x AZ21603/rev  ..
seq_name: gb_gss:AZ998589

seq_documentation_block:
LOCUS          54 bp      DNA
DEFINITION     clone UGGC2M0285D08 R, DNA sequence.
ACCESSION     AZ998589
VERSION       AZ998589.1  GI:13869816
KEYWORDS      GSS.
SOURCE        Mus musculus
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciuromorpha; Muridae; Murinae; Mus.
REFERENCE     1 (bases 1 to 54)
AUTHORS       Dunn,D., Avagci,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenin,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
TITLE         Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
UNPUBLISHED   Unpublished (2000)
CONTACT       Contact: Robert B. Weiss
JOURNAL      University of Utah Genome Center
COMMENT      Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
              Tel: 801 585 5606
              Fax: 801 585 7177
              Email: ddunne@genetics.utah.edu
PLATE         Insert Length: 10000 Std Error: 0.00
PLATE         Plate: 0285 Row: D Column: 08
SEQ_PRIMER    Seq primer: CACACAGGAAACAGCTATGACC
CLASS         Class: Plasmid ends
HIGH_QUALITY  High quality sequence stop: 54.
LOCATION      Location/Qualifiers
              1..54
ORGANISM     /organism="Mus musculus"

alignment_scores:
    Quality: 39.00      Length: 11
    Ratio: 4.333        Gaps: 0
Percent Similarity: 81.818      Percent Identity: 54.545
alignment_block:
Align seg 1/1 to reverse of: AZ21603  from: 1 to: 57
US-09-439-311-2 x AZ21603/rev  ..
seq_name: gb_gss:AZ998589

seq_documentation_block:
LOCUS          54 bp      DNA
DEFINITION     clone UGGC2M0285D08 R, DNA sequence.
ACCESSION     AZ998589
VERSION       AZ998589.1  GI:13869816
KEYWORDS      GSS.
SOURCE        Mus musculus
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciuromorpha; Muridae; Murinae; Mus.
REFERENCE     1 (bases 1 to 54)
AUTHORS       Dunn,D., Avagci,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmoud,M., Meenin,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A. and Wright,D., Weiss,R.
TITLE         Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
UNPUBLISHED   Unpublished (2000)
CONTACT       Contact: Robert B. Weiss
JOURNAL      University of Utah Genome Center
COMMENT      Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
              Tel: 801 585 5606
              Fax: 801 585 7177
              Email: ddunne@genetics.utah.edu
PLATE         Insert Length: 10000 Std Error: 0.00
PLATE         Plate: 0285 Row: D Column: 08
SEQ_PRIMER    Seq primer: CACACAGGAAACAGCTATGACC
CLASS         Class: Plasmid ends
HIGH_QUALITY  High quality sequence stop: 54.
LOCATION      Location/Qualifiers
              1..54
ORGANISM     /organism="Mus musculus"

/stain="C57BL/6J"
/db_xref="taxon:10090"
/clone_1ib="006 - Rescuemu Grid G"
/clone_2ib="Mouse 10kb plasmid UGGC2M library"
/sex="Female"
/lab_host="E. coli strain XL10-Gold, Tr-resitant, F-"
/note="Vector: PMW2nv; Purified genomic DNA from M. musculus C57BL/6J (female) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA Polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (91473214|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptored mouse DNA was annealed to adaptored vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance." (Stratagene) cells
BASE COUNT          11 a   2 c   21 g   20 t
ORIGIN

alignment_scores:
    Quality: 38.50      Length: 18
    Ratio: 2.750        Gaps: 1
Percent Similarity: 77.778      Percent Identity: 50.000
alignment_block:
Align seg 1/1 to: AZ998589  from: 1 to: 54
US-09-439-311-2 x AZ998589  ..
seq_name: gb_estl:AU106648

seq_documentation_block:
LOCUS          50 bp      mRNA
DEFINITION     Sugano Homo sapiens cDNA library Homo sapiens cDNA clone KAT05523, mRNA sequence.
ACCESSION     AU106648
VERSION       AU106648.1  GI:15556169
KEYWORDS      EST.
SOURCE        human.
ORGANISM      Homo sapiens
REFERENCE     1 (bases 1 to 50)
AUTHORS       Suzuki,Y., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata,H., Ota,T., Isouai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo,K., Sugama,A. and Sugano,S.
TITLE         Fine structural analysis of transcription start sites of human mRNAs using full-length enriched and 5'-end enriched cDNA libraries
UNPUBLISHED   Unpublished (2001)
COMMENT      Contact: Yuraku Suzuki
              Department of Virology
              Institute of Medical Science, University of Tokyo
              4-6-1, Shirokanedai, Minato-ku, Tokyo 108-8639, Japan
              Email: ysuzuki@ims.u-tokyo.ac.jp
FEATURES      source
              /organism="Mus musculus"

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seq\_name: gb-est1:AU102739

seq\_documentation\_block:

LOCUS AU102739 mRNA Homo sapiens cDNA library Homo sapiens CDNA clone

DEFINITION HRC13119, mRNA sequence.

ACCESSION AU102739

VERSION AU102739.1 GI:13552560

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.

REFERENCE 1 (bases 1 to 50)

AUTHORS Suzuki,I., Tsunoda,T., Taira,H., Mizushima-Sugano,J., Sese,J., Hata ,H., Ota,T., Isogai,T., Tanaka,T., Nakamura,Y., Morishita,S., Okubo K., Sugama,A. and Sugano,S.

TITLE Fine structural analysis of transcription start sites of human mRNAs using full-length enriched and 5'-end enriched cDNA libraries

JOURNAL Unpublished (2001)

COMMENT Contact: Yutaka Suzuki

Institute of Medical Science, University of Tokyo  
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan  
Email: ysuuki@ims.u-tokyo.ac.jp

Suzuki,Y., Yoshitomo,Nakada,K., Maruyama,K., Sugama,A. and Sugano ,S. Construction and characterization of a full length-enriched and a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES

source

1. .50

/organism="Homo sapiens"

/db\_xref="txon:9606"

/clone="HRC13119"

/clone.lib="Sugano Homo sapiens cDNA library"

BASE COUNT

7 a 16 c 18 g 9 t

ORIGIN

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Ratio:	2.846		
Percent Identity:	92.857		

alignment\_block:  
US-09-439-311-2 x AU102739/rev

Align seg 1/1 to reverse of: AU102739 from: 1 to: 50

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||||::||||::|||:|||:|||:|||:|||:|||:|||:|||:|||:|||:  
42 ACTAGGGCCCCGCCTACAGGGAGCTCCATTCTCCGCACG 1